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(54) Title: 70 HUMAN SECRETED PROTEINS			
(57) Abstract			
<p>The present invention relates to 70 novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins.</p>			

60/043,314	11 April 1997 (11.04.97)	US	60/047,598	23 May 1997 (23.05.97)	US	60/056,882	22 August 1997 (22.08.97)	US
60/043,569	11 April 1997 (11.04.97)	US	60/047,613	23 May 1997 (23.05.97)	US	60/056,637	22 August 1997 (22.08.97)	US
60/043,311	11 April 1997 (11.04.97)	US	60/047,582	23 May 1997 (23.05.97)	US	60/056,903	22 August 1997 (22.08.97)	US
60/043,671	11 April 1997 (11.04.97)	US	60/047,596	23 May 1997 (23.05.97)	US	60/056,888	22 August 1997 (22.08.97)	US
60/043,674	11 April 1997 (11.04.97)	US	60/047,612	23 May 1997 (23.05.97)	US	60/056,879	22 August 1997 (22.08.97)	US
60/043,669	11 April 1997 (11.04.97)	US	60/047,632	23 May 1997 (23.05.97)	US	60/056,880	22 August 1997 (22.08.97)	US
60/043,312	11 April 1997 (11.04.97)	US	60/047,601	23 May 1997 (23.05.97)	US	60/056,894	22 August 1997 (22.08.97)	US
60/043,313	11 April 1997 (11.04.97)	US	60/047,595	23 May 1997 (23.05.97)	US	60/056,911	22 August 1997 (22.08.97)	US
60/043,672	11 April 1997 (11.04.97)	US	60/047,599	23 May 1997 (23.05.97)	US	60/056,636	22 August 1997 (22.08.97)	US
60/043,315	11 April 1997 (11.04.97)	US	60/047,588	23 May 1997 (23.05.97)	US	60/056,874	22 August 1997 (22.08.97)	US
60/043,578	11 April 1997 (11.04.97)	US	60/047,585	23 May 1997 (23.05.97)	US	60/056,910	22 August 1997 (22.08.97)	US
60/043,576	11 April 1997 (11.04.97)	US	60/047,586	23 May 1997 (23.05.97)	US	60/056,864	22 August 1997 (22.08.97)	US
60/043,670	11 April 1997 (11.04.97)	US	60/047,590	23 May 1997 (23.05.97)	US	60/056,631	22 August 1997 (22.08.97)	US
60/047,600	23 May 1997 (23.05.97)	US	60/047,594	23 May 1997 (23.05.97)	US	60/056,845	22 August 1997 (22.08.97)	US
60/047,615	23 May 1997 (23.05.97)	US	60/047,589	23 May 1997 (23.05.97)	US	60/056,892	22 August 1997 (22.08.97)	US
60/047,597	23 May 1997 (23.05.97)	US	60/047,593	23 May 1997 (23.05.97)	US	60/056,632	22 August 1997 (22.08.97)	US
60/047,502	23 May 1997 (23.05.97)	US	60/047,614	23 May 1997 (23.05.97)	US	60/056,664	22 August 1997 (22.08.97)	US
60/047,633	23 May 1997 (23.05.97)	US	60/047,501	23 May 1997 (23.05.97)	US	60/056,876	22 August 1997 (22.08.97)	US
60/047,583	23 May 1997 (23.05.97)	US	60/048,974	06 June 1997 (06.06.97)	US	60/056,881	22 August 1997 (22.08.97)	US
60/047,617	23 May 1997 (23.05.97)	US	60/048,964	06 June 1997 (06.06.97)	US	60/056,909	22 August 1997 (22.08.97)	US
60/047,618	23 May 1997 (23.05.97)	US	60/056,886	22 August 1997 (22.08.97)	US	60/056,875	22 August 1997 (22.08.97)	US
60/047,503	23 May 1997 (23.05.97)	US	60/056,877	22 August 1997 (22.08.97)	US	60/056,862	22 August 1997 (22.08.97)	US
60/047,592	23 May 1997 (23.05.97)	US	60/056,889	22 August 1997 (22.08.97)	US	60/056,887	22 August 1997 (22.08.97)	US
60/047,581	23 May 1997 (23.05.97)	US	60/056,893	22 August 1997 (22.08.97)	US	60/056,908	22 August 1997 (22.08.97)	US
60/047,584	23 May 1997 (23.05.97)	US	60/056,630	22 August 1997 (22.08.97)	US	60/056,884	22 August 1997 (22.08.97)	US
60/047,500	23 May 1997 (23.05.97)	US	60/056,878	22 August 1997 (22.08.97)	US	60/057,761	05 September 1997 (05.09.97)	US
60/047,587	23 May 1997 (23.05.97)	US	60/056,662	22 August 1997 (22.08.97)	US	60/057,650	05 September 1997 (05.09.97)	US
60/047,492	23 May 1997 (23.05.97)	US	60/056,872	22 August 1997 (22.08.97)	US			

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70 Human Secreted Proteins

Field of the Invention

This invention relates to newly identified polynucleotides and the polypeptides encoded by these polynucleotides, uses of such polynucleotides and polypeptides, and
5 their production.

Background of the Invention

Unlike bacterium, which exist as a single compartment surrounded by a membrane, human cells and other eucaryotes are subdivided by membranes into many functionally distinct compartments. Each membrane-bounded compartment, or
10 organelle, contains different proteins essential for the function of the organelle. The cell uses "sorting signals," which are amino acid motifs located within the protein, to target proteins to particular cellular organelles.

One type of sorting signal, called a signal sequence, a signal peptide, or a leader sequence, directs a class of proteins to an organelle called the endoplasmic reticulum
15 (ER). The ER separates the membrane-bounded proteins from all other types of proteins. Once localized to the ER, both groups of proteins can be further directed to another organelle called the Golgi apparatus. Here, the Golgi distributes the proteins to vesicles, including secretory vesicles, the cell membrane, lysosomes, and the other organelles.

20 Proteins targeted to the ER by a signal sequence can be released into the extracellular space as a secreted protein. For example, vesicles containing secreted proteins can fuse with the cell membrane and release their contents into the extracellular space - a process called exocytosis. Exocytosis can occur constitutively or after receipt of a triggering signal. In the latter case, the proteins are stored in secretory vesicles (or
25 secretory granules) until exocytosis is triggered. Similarly, proteins residing on the cell membrane can also be secreted into the extracellular space by proteolytic cleavage of a "linker" holding the protein to the membrane.

Despite the great progress made in recent years, only a small number of genes encoding human secreted proteins have been identified. These secreted proteins include
30 the commercially valuable human insulin, interferon, Factor VIII, human growth hormone, tissue plasminogen activator, and erythropoietin. Thus, in light of the pervasive role of secreted proteins in human physiology, a need exists for identifying and characterizing novel human secreted proteins and the genes that encode them. This knowledge will allow one to detect, to treat, and to prevent medical disorders by using
35 secreted proteins or the genes that encode them.

Summary of the Invention

The present invention relates to novel polynucleotides and the encoded polypeptides. Moreover, the present invention relates to vectors, host cells, antibodies, and recombinant methods for producing the polypeptides and polynucleotides. Also provided are diagnostic methods for detecting disorders related to the polypeptides, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying binding partners of the polypeptides.

Detailed Description

Definitions

The following definitions are provided to facilitate understanding of certain terms used throughout this specification.

In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide.

In the present invention, a "secreted" protein refers to those proteins capable of being directed to the ER, secretory vesicles, or the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily containing a signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage.

As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence contained in SEQ ID NO:X or the cDNA contained within the clone deposited with the ATCC. For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, with or without the signal sequence, the secreted protein coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having the translated amino acid sequence generated from the polynucleotide as broadly defined.

In the present invention, the full length sequence identified as SEQ ID NO:X was often generated by overlapping sequences contained in multiple clones (contig

analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X was deposited with the American Type Culture Collection ("ATCC"). As shown in Table 1, each clone is identified by a cDNA Clone ID (Identifier) and the ATCC Deposit Number. The ATCC is located at 12301 Park Lawn Drive, Rockville, Maryland 20852, USA. The ATCC deposit was made pursuant to the terms of the Budapest Treaty on the international recognition of the deposit of microorganisms for purposes of patent procedure.

A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, the complement thereof, or the cDNA within the clone deposited with the ATCC. "Stringent hybridization conditions" refers to an overnight incubation at 42°C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH_2PO_4 ; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 µg/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a

complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

5 The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and
10 double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability
15 or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine. A variety of modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.

 The polypeptide of the present invention can be composed of amino acids joined
20 to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs,
25 as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be
30 branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a
35 nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine,

formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, **PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES**, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); **POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS**, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., *Meth Enzymol* 182:626-646 (1990); Rattan et al., *Ann NY Acad Sci* 663:48-62 (1992).)

"SEQ ID NO:X" refers to a polynucleotide sequence while "SEQ ID NO:Y" refers to a polypeptide sequence, both sequences identified by an integer specified in Table 1.

"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.)

Polynucleotides and Polypeptides of the Invention

FEATURES OF PROTEIN ENCODED BY GENE NO: 1

The translation product of Gene NO: 1 shares sequence homology with alpha-L-fucosidase which is thought to be important as a lysosomal enzyme that hydrolyzes fucose from fucoglycoconjugates. (See Accession No. gi/178409.) Lysosome fructosidase is involved in certain lysosome storage diseases. (See *Biochem. Biophys. Res. Commun.*, 164(1):439-445 (1989).) Fucosidosis, an autosomal recessive lysosomal storage disorder characterized by progressive neurological deterioration and mental retardation. The disease results from deficient activity of alpha-L-fucosidase, a lysosomal enzyme that hydrolyzes fucose from fucoglycoconjugates. This gene likely encodes a novel fucosidase isoenzyme. Based on homology, it is likely that the translated product of this gene is also involved in lysosome catabolism of molecules and

that aberrations in the concentration and/or composition of this product may be causative in lysosome storage disorders. Preferred polypeptide fragments comprise the amino acid sequence PGHLLPHKWENC (SEQ ID NO: 257).

Gene NO: 1 is expressed primarily in stromal cells, and to a lesser extent in
5 human fetal kidney and human tonsils.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, fucosidosis and other lysosome storage disorders. Similarly,
10 polypeptides and antibodies directed to the polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues of cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., stromal cells, kidney, tonsils, and cancerous and
15 wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology of Gene NO: 1 to alpha-L-fucosidase
20 indicates that polypeptides and polynucleotides corresponding to Gene NO: 1 are useful for the treatment of fucosidosis and general lysosomal disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 134 as residues: Met-1 to Leu-6, Thr-32 to Glu-39, Lys-80 to Lys-85, and Met-90 to
25 Pro-96.

FEATURES OF PROTEIN ENCODED BY GENE NO: 2

The translation product of Gene No. 2 shares sequence homology with stromal cell-derived factor-2 (SDF-2) which is a novel secreted factor. See, for example, Gene, 176(1-2):211-214, (1996, Oct. 17.) The amino acid sequence of SDF-2 shows
30 similarity to yeast dolichyl phosphate-D-mannose:protein mannosyltransferases, Pmt1p [Strahl-Bolsinger et al. Proc. Natl. Acad. Sci. USA 90, 8164-8168 (1993)] and Pmt2p [Lussier et al. J. Biol. Chem. 270, 2770-2775 (1995)], whose activities have not been detected in higher eukaryotes. Based on the sequence similarity, the translation product of this gene is expected to share certain biological activities with SDF-2, Pmt1p and
35 Pmt2p.

Gene NO: 2 is expressed primarily in immune system tissue and cancerous tissues, such as liver hepatoma, human B-cell lymphoma, spleen in a patient suffering

from chronic lymphocytic leukemia, hemangiopericytoma, pharynx carcinoma, breast cancer, thyroid, bone marrow, osteoblasts and to a lesser extent in a few other tissues such as kidney pyramids.

Therefore, polynucleotides or polypeptides of the invention are useful as
5 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of the diseases and conditions which include, but are not limited to, disorders in kidney, liver, and immune organs, particularly cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in
10 providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the kidney, liver, thyroid, and bone marrow expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., liver, spleen, B-cells, pharynx, thyroid, mammary tissue, bone marrow, osteoblasts and
15 kidneys, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology of Gene NO: 2 to stromal cell-derived
20 factor-2 indicates that polypeptides and polynucleotides corresponding to Gene NO: 2 are useful for diagnosis and therapeutic treatment of disorders in kidney, liver, and immune organs since stromal cells play important role in organ function. Stroma carries the blood supply and provides support for the growth of parenchymal cells and is therefore crucial to the growth of a neoplasm. Nucleic acids of the present invention
25 comprise, but preferably do not consist of, and more preferably do not comprise, SEQ ID NO: 3 from US Patent No. 5,576,423, incorporated herein by reference, and shown herein as SEQ ID NO: 258).

Preferred epitopes include those comprising a sequence shown in SEQ ID NO:
135 as residues: His-56 to Gly-65, Ala-74 to Ser-80, Ile-84 to Pro-97, Leu-124 to Glu-
30 129, Glu-135 to Asp-143, Gly-175 to Ser-180, and Ala-194 to Thr-199.

FEATURES OF PROTEIN ENCODED BY GENE NO: 3

The translation product of Gene NO: 3 shares sequence homology with LZIP-1, LZIP-2 and other leucine zipper proteins, which are thought to be important in nucleic
35 acid binding. This gene has been reported in Mol. Cell. Biol. 17 (9), 5117-5126 (1997) as "Luman". Luman is a cyclic AMP response element (CRE)-binding protein/activating transcription factor 1 protein of the basic leucine zipper superfamily. It binds CREs in

vitro and activates CRE-containing promoters when transfected into COS7 cells. The complete amino acid sequence of Luman reported in Mol. Cell. Biol. 17 (9): 5117-5126 (1997) is:

MELELDAGDQDLLAFLLEESGDLGTAPDEAVRAPLDWALPLSEVPSDWEVDDLL
 5 CSLLSPPASLNILSSSNPCLVHHHTYSLPRETVSMDLESESCRKEGTQMTPQH
 MEELAEQEIARLVLTDEEKSLLEKEGLILPETLPLTKTEEQILKRVRKIRNKRSA
 QESRRKKKVYVGGLESRLKYTAQNMEQLQNKVQLLEEQNLSLLDQLRKLQAM
 VIEISNKTSSSSTCILVLLVSFCLLLVPAMYSSDTRGSLPAEHGVLSRQLRALPSE
 DPYQLELPALQSEVPKDSHTQWLDGSDCVLQAPGNTSCLLHYMPQAPSAEPPL
 10 EWPFPDLSS EPLCRGPILPLQANLTRKGGWLPTGSPSVILQDRYSG (SEQ ID
 N:259).

Gene NO: 3 is expressed primarily in apoptotic T-cells and Soares senescent cells and to a lesser extent in multiple tissues and cell types, including, multiple sclerosis tissue, and hippocampus.

15 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, immunologically mediated disorders, transplantation, immunodeficiency, and tumor necrosis. Similarly, polypeptides and antibodies directed to these
 20 polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and transplantation, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., multiple sclerosis tissue, hippocampus, bone marrow and cancerous and wounded
 25 tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology of Gene NO: 3 to leucine zipper nucleic
 30 acid binding proteins indicates that polypeptides and polynucleotides corresponding to Gene NO: 3 are useful for diagnosis and treatment of immunologically mediated disorders, transplantation, immunodeficiency, and tumor necrosis. The secreted nucleic acid binding protein in the apoptotic tissues may be involved in the disposal of the DNA released by apoptotic cells. Furthermore, the studies conducted in support of Luman
 35 suggest that the translation product of this gene may be used to identify transcriptional regulation elements which in turn are useful in modulation of immune function.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 136 as residues: Asn-7 to Ser-12, Tyr-32 to Gly-38, Pro-55 to Tyr-60, Glu-70 to Thr-76, and Pro-104 to Leu-110.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 4

The translation product of Gene NO: 4 shares sequence homology with a number of tetraspan transmembrane surface molecules such as human metastasis tumor suppressor gene, CO-029 tumor associated antigen protein, CD53 hematopoietic antigen, human membrane antigen TM4 superfamily protein, metastasis controlling
10 peptide, and human CD9 sequence, which are thought to be important in development of cancer, immune system development and functions.

Gene NO: 4 is expressed primarily in cancers of several different tissues and to a lesser extent in normal tissue like prostate, skin and kidney.

Therefore, polynucleotides or polypeptides of the invention are useful as
15 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancers and disorders of the immune system, prostate and kidney. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell
20 type(s). For a number of disorders of the above tissues or cells, particularly of the kidney, skin, prostate and immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., kidney, skin and prostate, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from
25 an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology of Gene NO: 4 to tetraspan transmembrane surface molecules such as human metastasis tumor suppressor gene,
30 CO-029 tumor associated antigen protein, CD53 hematopoietic antigen, human membrane antigen TM4 superfamily protein, metastasis controlling peptide, and human CD9 sequence, indicates that polypeptides and polynucleotides corresponding to Gene NO: 4 are involved with the cellular control of growth and differentiation. Therefore, the translation product of this gene is believed to be useful for diagnosis and treatment
35 of neoplasia and disorders of the kidney, skin and prostate. For example, recombinant protein can be produced in transformed host cells for diagnostic and prognostic applications. Alterations in the protein sequence are indicative of the presence of

malignant cancer, or of a predisposition to malignancy, in a subject. Gene therapy can be used to restore the wild-type gene product to a subject. Additionally, the antibodies are a useful tool for the identification of hematopoietic neoplasms, and may prove helpful for identifying morphologically poorly defined cells. Moreover, this protein can be used to isolate cognate receptors and ligands and identify potential agonists and antagonists using techniques known in the art. The protein also has immunomodulatory activity, regulates hematopoiesis and stimulates growth and regeneration as a male/female contraceptive, increases fertility depending on activin and inhibin like activities. Other uses are as a chemotactic agent for lymphocytes, treatment of coagulation disorders, an anti-inflammatory agent, an antimicrobial or analgesic and as a modulator of behavior and metabolism. The DNA can be used in genetic diagnosis or gene therapy, and for the production of recombinant protein. It can also be used to identify protein expressing cells, isolate related sequences, prepare primers for genetic fingerprinting and generate anti-protein or anti-DNA antibodies. In addition, residues 1-71, in the translation product for this gene are believed to be the extracellular domain. Thus, polypeptide comprising residues 1-71 or derivatives (including fragments) or analogs thereof, are useful as a soluble polypeptide which may be routinely used therapeutically to antagonize the activities of the receptor.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 137 as residues: Lys-118 to Phe-127, Asn-145 to Ala-160, and Thr-177 to Val-188.

FEATURES OF PROTEIN ENCODED BY GENE NO: 5

Gene NO: 5 is expressed primarily in human testes.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, diseases of the testes including cancer and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., testes and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of Gene NO: 5 indicates that the protein product of this gene is useful for treatment/diagnosis of diseases of the testes, particularly testicular cancer since expression is observed primarily in the testes.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO:
5 138 as residue: Gly-22 to Gln-30.

FEATURES OF PROTEIN ENCODED BY GENE NO: 6

The translation product of Gene NO: 6 shares sequence homology with GALNS (N-acetylgalactosamine 6-sulphatase) which is thought to be important in the storage of
10 the glycosaminoglycans, keratan sulfate and chondroitin 6-sulfate. See Genbank accession no. gil618426. Based on the sequence similarity, the translation product of this gene is expected to share biological activities with GALNS.

Gene NO: 6 is expressed primarily in human bone marrow.

Therefore, polynucleotides or polypeptides of the invention are useful as
15 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, storage disorders of glycosaminoglycans, keratan sulfate and chondroitin 6-sulfate, e.g., Morquio A syndrome. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential
20 identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly involving cell storage disorder, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., bone marrow and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from
25 an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology of Gene NO: 6 to N-acetylgalactosamine 6-sulphatase indicates that polypeptides and polynucleotides corresponding to Gene
30 NO: 6 are useful for the treatment and diagnosis of storage disorders of glycosaminoglycans, keratan sulfate and chondroitin 6-sulfate. Such disorders are known in the art and include, e.g., Morquio A syndrome which is caused by an error of mucopolysaccharide metabolism with excretion of keratan sulfate in urine. Morquio A syndrome is characterized by severe skeletal defects with short stature, severe deformity
35 of spine and thorax, long bones with irregular epiphyses but with shafts of normal length, enlarged joints, flaccid ligaments, and waddling gait: autosomal recessive inheritance.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 139 as residues: Gly-29 to Pro-36 and Glu-57 to Leu-64.

FEATURES OF PROTEIN ENCODED BY GENE NO: 7

5 The translation product of Gene NO: 7 shares sequence homology with carboxy peptidase E and H (carboxypeptidase E is thought to be important in the biosynthesis of numerous peptide hormones and neurotransmitters). The translation product of this gene also shares sequence homology with bone-related carboxypeptidase "OSF-5" from the mouse. See European patent application EP-588118-A. Based on the sequence
10 similarity to OSF-5, the translation product of this gene will hereinafter sometimes be referred to as "human-OSF-5" or "hOSF-5".

Gene NO: 7 is expressed primarily in tumor cell lines derived from connective tissues including chondrosarcoma, synovial sarcoma, Wilm's tumor and rhabdomyosarcoma and to a lesser extent in a myeloid progenitor cell line, bone
15 marrow, and placenta.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, various cancers involving the skeletal system and connective tissues in
20 general, in particular at cartilage interfaces. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal system and various other tumor tissues, expression of this gene at significantly higher or lower levels may routinely be
25 detected in certain tissues (e.g., connective tissues and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

30 The restricted tissue distribution and homology of Gene NO: 7 to carboxypeptidase E and mouse OSF-5 indicates that polypeptides and polynucleotides corresponding to Gene NO: 7 are for processing of peptides to their mature form that may have various activities similar to the activities of neuropeptides but in the periphery. In addition the abundance of expression in cancer tissues indicates that
35 aberrant expression and subsequent processing may play a role in the progression of malignancies, e.g., growth factor and/or adhesion factor activities. In particular, the expression of this gene is restricted to connective tissues and embryonic tissues.

Furthermore, it is overexpressed in cancers of these same tissues (i.e., in sarcomas). Moreover, hOSF-5 shares very strong sequence similarity with mOSF-5 which is a known bone growth factor and is thought to be useful in obtaining products for the diagnosis and treatment of bone metabolic diseases, e.g., osteoporosis and Paget's disease. Like OSF-5, the translation product of this gene is believed to be a bone-specific carboxypeptidase which acts as an adhesion molecule/growth factor and takes part in osteogenesis at the site of bone induction. hOSF-5 can, therefore, be used to treat bone metabolic diseases, osteoporosis, Paget's disease, osteomalacia, hyperostosis or osteopetrosis. Furthermore, hOSF-5 can be used to stimulate the regeneration of bone at the site of mechanical damage, e.g., accidentally or surgically caused fractures.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 140 as residues: Leu-24 to Val-30, Ala-89 to Lys-94, Phe-150 to Trp-157, Leu-162 to Asp-167, Asp-187 to Ser-199, His-241 to Asp-254, and Pro-362 to Asp-376.

15

FEATURES OF PROTEIN ENCODED BY GENE NO: 8

Gene NO: 8 is expressed primarily in bone marrow, and to a lesser extent in an erythroleukemia cell line.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, hematological disorders including cancer and anemia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and hematologic systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., bone marrow, kidney, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 8 are useful as a growth factor for hematopoietic stem cells or progenitor cells, e.g., in the treatment of bone marrow stem cell loss in chemotherapy patients and in the treatment of kidney disease.

35

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 141 as residues: Gly-30 to Lys-35.

FEATURES OF PROTEIN ENCODED BY GENE NO: 9

Gene NO: 9 is expressed primarily in neutrophils.

Therefore, polynucleotides or polypeptides of the invention are useful as
5 reagents for differential identification of the cell type present in a biological sample and
for diagnosis of diseases and conditions which include, but are not limited to,
inflammatory diseases. Similarly, polypeptides and antibodies directed to these
polypeptides are useful in providing immunological probes for differential identification
10 of the cell type indicated. For a number of disorders of the above tissues or cells,
particularly of the immune system, expression of this gene at significantly higher or
lower levels may routinely be detected in certain tissues or cell types (e.g., neutrophils,
bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum,
plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from
15 an individual having such a disorder, relative to the standard gene expression level, i.e.,
the expression level in healthy tissue or bodily fluid from an individual not having the
disorder.

The tissue distribution indicates that polypeptides and polynucleotides
corresponding to Gene NO: 9 are useful for immune modulation or as a growth factor
to stimulate neutrophil differentiation or proliferation that may be useful in the treatment
20 of neutropenia.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO:
142 as residues: Thr-22 to Pro-37.

FEATURES OF PROTEIN ENCODED BY GENE NO: 10

25 Gene NO: 10 is expressed primarily in the epidermis.

Therefore, polynucleotides or polypeptides of the invention are useful as
reagents for differential identification of the tissue(s) or cell type(s) present in a
biological sample and for diagnosis of diseases and conditions which include, but are
not limited to, diseases of the epidermis such as psoriasis or eczema or may be involved
30 in the normal proliferation or differentiation of the epithelial cells or fibroblasts
constituting the skin. Similarly, polypeptides and antibodies directed to these
polypeptides are useful in providing immunological probes for differential identification
of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,
particularly of the skin, expression of this gene at significantly higher or lower levels
35 may routinely be detected in certain tissues (e.g., epidermis and cancerous and
wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal
fluid) or another tissue or cell sample taken from an individual having such a disorder.

relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 10 are useful for diagnosis and treatment of skin conditions and as an aid in the healing of various epidermal injuries including wounds, and diabetic ulcers.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 143 as residues: Ser-3 to Ser-9 and Trp-27 to Glu-32.

10 FEATURES OF PROTEIN ENCODED BY GENE NO: 11

The translation product of Gene NO: 11 shares sequence homology with phosphatidylcholine 2-acylhydrolase (PLA2). See, for example, Genbank accession no. gil190004. PLA2 is involved in inflammation, where it is responsible for the conversion of cell membrane phospholipids into arachidonic acid. Arachidonic acid in turn feeds into both the lipoxygenase and cyclooxygenase pathways to produce leukotrienes (involved in chemotaxis, vasoconstriction, bronchoconstriction, and increased vascular permeability) and prostaglandins (responsible for vasodilation, potentiate edema, and increased pain). Diseases in which PLA2 is implicated as a major factor include rheumatoid arthritis, sepsis, ischemia, and thrombosis. The inventors refer to the translation product of this gene as PLA2-like protein based on the sequence similarity. Furthermore, owing to the sequence similarity PLA2 and PLA2-like protein are expected to share certain biological activities.

Gene NO: 11 is expressed primarily in human cerebellum and in T-cells.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cerebellum disorders, rheumatoid arthritis, sepsis, ischemia, and thrombosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cerebellum and Purkinje cells, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., brain, bone marrow, T-cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 11 are useful for diagnosis and treatment of cerebellum disorders, rheumatoid arthritis, sepsis, ischemia, and thrombosis. This gene is also useful as a chromosome marker. It is believed to map to Chr.15, D15S118-D15S123.

5

FEATURES OF PROTEIN ENCODED BY GENE NO: 12

Gene NO: 12 is expressed primarily in highly vascularized tissues such as placenta, uterus, tumors, fetal liver, fetal spleen and also in the C7MCF7 cell line treated with estrogen.

10

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, endometriosis, endometritis, endometrial carcinoma, primary hepatocellular carcinoma, and spleen-related diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endometrium, liver and spleen, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., endometrium, liver, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

15

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The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 12 are useful for diagnosis and treatment of diseases of the endometrium (such as endometrial carcinoma, endometriosis, and endometritis), liver diseases (such as primary hepatocellular carcinoma), and spleen-related diseases.

SEQ ID NO: 145 as residues: Ala-29 to Leu-35, Leu-50 to Ser-57, Glu-96 to Glu-105, Asp-140 to Asp-148, and Asn-191 to Ser-197.

30

FEATURES OF PROTEIN ENCODED BY GENE NO: 13

Gene NO: 13 is expressed primarily in B cell lymphoma and to a lesser extent in other tissues.

35

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, B cell lymphoma; hematopoietic disorders; immune dysfunction.

Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may
5 routinely be detected in certain tissues and cell types (e.g., bone marrow and B-cells and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the
10 disorder.

Enhanced expression of this gene product in B cell lymphoma indicates that it may play a role in the proliferation of hematopoietic cells. It is also believed to be involved in the survival and/or differentiation of various hematopoietic lineages. Expression in lymphoma also indicates that it may be involved in other cancers and
15 abnormal cellular proliferation. The tissue distribution, therefore, indicates that polypeptides and polynucleotides corresponding to Gene NO: 13 are useful for the diagnosis and/or therapeutic treatment of hematopoietic disorders, particularly B cell lymphoma. Furthermore, since overexpression of this gene is associated with the development of B cell lymphoma, antagonists of this protein are useful to interfere with
20 the progression of the disease. This protein is useful in assays for identifying such antagonists. Assays for identifying antagonists are known in the art and are described briefly elsewhere herein. Preferred antagonists include antibodies and antisense nucleic acid molecules. Preferred are antagonists which inhibit B-cell proliferation.

25 **FEATURES OF PROTEIN ENCODED BY GENE NO: 14**

The translation product of Gene NO: 14 shares sequence homology with very low density lipoprotein receptor which is thought to be important in transport of lipoproteins. Owing to the sequence similarity the translation product of this gene is believed to share certain biological activities with VLDL receptors. Assaying such
30 activity may be achieved by assays known in the art and set forth elsewhere herein.

This gene is expressed primarily in human synovium, umbilical vein endothelial cells, CD34+ cells, Jurkat cells, and HL60 cells, and to a lesser extent in thymus, meningioma, hypothalamus, adult testis, and fetal liver and spleen.

Therefore, polynucleotides or polypeptides of the invention are useful as
35 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, atherosclerosis, ataxia malabsorption, vascular damage, hyperlipidemia,

and other cardiovascular diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular and hematological systems,
5 expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., endothelium, thymus meningioma, hypothalamus, testes, liver, and spleen and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,
10 the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in the vascular endothelial cells and homology to VLDL receptors indicates that polypeptides and polynucleotides corresponding to Gene NO: 14 are useful for diagnosis and treatment of atherosclerosis, ataxia malabsorption, and
15 hyperlipidemia. These and other factors often result in other cardiovascular diseases. Additionally, the presence of the gene product in cells of blood lineages indicates that it may be useful in hematopoietic regulation and hemostasis.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 147 as residues: Pro-39 to Ser-52, Trp-71 to Thr-76, and Pro-94 to His-100.

20

FEATURES OF PROTEIN ENCODED BY GENE NO: 15

The translation product of Gene NO: 15 shares sequence homology with kallikrein which is thought to be important in blood pressure and renal secretion. Furthermore, this gene has now been characterized as a novel hepatitis B virus X
25 binding protein that inhibits viral replication. See, for example, J. Virol. 72 (3), 1737-1743 (1998).

This gene is expressed primarily in kidney, placenta, lung, aorta and other endothelial cells, caudate nucleus and to a lesser extent in melanocytes, liver, adipose tissue.

30 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, renovascular hypertension, renal secretion, electrolyte metabolism, toxemia of pregnancy. Similarly, polypeptides and antibodies directed to these
35 polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the renovascular or respiratory vascular systems, expression of this gene

at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., kidney, placenta, lung, endothelial cells, melanocytes, liver, and adipose tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to kallikrein indicates that polypeptides and polynucleotides corresponding to Gene NO: 15 are useful for treating renovascular hypertension, renal secretion, electrolyte metabolism, toxemia of pregnancy and hydronephrosis. The protein expression in the organs like kidney, lung and vascular endothelial cells indicates the gene involvement in hemodynamic regulatory functions. The translation product of this gene is also useful in the treatment of viral infection, particularly liver infection, and particularly hepatitis B virus(es).

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 148 as residues: Leu-9 to Asn-15 and Thr-56 to Asp-61.

FEATURES OF PROTEIN ENCODED BY GENE NO: 16

The translation product of Gene NO: 16 shares sequence homology with secretory component protein, immunoglobulins and their receptors which are thought to be important in immunological functions. The amino acid sequence of secretory component protein can be accessed as accession no. pirlA02112, incorporated herein by reference.

Gene NO: 16 is expressed primarily in macrophages, monocytes and dendritic cells and to a lesser extent in placenta and brain.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, inflammation and tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues or cells (e.g., macrophages, monocytes, dendritic cells, placenta and brain, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a

disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to immunoglobulins and secretory component protein indicates that polypeptides and polynucleotides corresponding to
5 Gene NO: 16 are useful for diagnosis and treatment of inflammation and bacterial infection, and other diseases where immunomodulation would be beneficial.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 149 as residues: Pro-37 to Cys-51, Gln-53 to Cys-60, Asn-99 to Gly-106, Gly-145 to Glu-151, and Ile-159 to Ser-164.

10

FEATURES OF PROTEIN ENCODED BY GENE NO: 17

The translation product of Gene NO: 17 is evolutionarily conserved and shares sequence homology with proteins from yeast and *C. elegans*. See, for example, Genbank accession no.gil746540. As is known in the art, strong sequence similarity to
15 a secreted protein from *C. elegans* is predictive of cellular location of human proteins.

Gene NO: 17 is expressed primarily in colon carcinoma cell lines, messangial cells, many tumors like T cell lymphoma, osteoclastoma, Wilm's tumor, adrenal gland tumor, testes tumor, synovial sarcoma, and to a lesser extent in placenta, lung and brain.

20

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, rapidly growing/dividing cells such as cancerous tissue, including, colon carcinoma, lymphomas, and sarcomas. Similarly, polypeptides and antibodies directed
25 to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the gastrointestinal, hematological and immune systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., placenta, lung, brain, colon, messangial cells, adrenal gland, T-cells, testes, and lymph tissue, and cancerous and wounded tissues) or
30 bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

35

The tissue distribution in colon cancer and many other tumors indicates that the polynucleotides and polypeptides of Gene NO: 17 are useful for cancer diagnosis and therapeutic targeting. The extracellular nature may contribute to solid tumor

immunosuppression, angiogenesis and cell growth stimulation. The tissue distribution of this gene in cells of the immune system indicates that polypeptides and polynucleotides corresponding to Gene NO: 17 are useful for treatment, prophylaxis and diagnosis of immune and autoimmune diseases, such as lupus, transplant rejection, allergic reactions, arthritis, asthma, immunodeficiency diseases, leukemia, and AIDS. Its expression predominantly in hematopoietic cells also indicates that the gene could be important for the treatment and/or detection of hematopoietic disorders such as graft versus host reaction, graft versus host disease, transplant rejection, myelogenous leukemia, bone marrow fibrosis, and myeloproliferative disease. The protein can also be used to enhance or protect proliferation, differentiation and functional activation of hematopoietic progenitor cells such as bone marrow cells, which could be useful for cancer patients undergoing chemotherapy or patients undergoing bone marrow transplantation. The protein may also be useful to increase the proliferation of peripheral blood leukocytes, which could be useful in the combat of a range of hematopoietic disorders including immunodeficiency diseases, leukemia, and septicemia.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 150 as residues: Val-131 to Asn-136.

FEATURES OF PROTEIN ENCODED BY GENE NO: 18

The translation product of Gene NO: 18 shares sequence homology with immunoglobulin, which is thought to be important in immunoreactions.

Gene NO: 18 is expressed primarily in macrophage.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., macrophage and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in macrophages and the weak homology to immunoglobulin indicates that polypeptides and polynucleotides corresponding to Gene

NO: 18 are useful for diagnosing and treating immune response disorders, including inflammation, antigen presentation and immunosurveillance.

FEATURES OF PROTEIN ENCODED BY GENE NO: 19

5 The translation product of Gene NO: 19 shares sequence homology with proline rich proteins which are thought to be important in protein-protein interaction.

 This gene has a wide range of tissue distribution, but is expressed primarily in normal prostate, synovial fibroblasts, brain amygdala depression, fetal bone and fetal cochlea, and to a lesser extent in adult retina, umbilical vein endothelial cells, atrophic
10 endometrium, osteoclastoma, melanocytes, pancreatic carcinoma and smooth muscle.

 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer metastasis, wound healing, tissue repair. Similarly, polypeptides
15 and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal, connective tissues, reproductive and central nervous system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., brain,
20 prostrate, fibroblasts, bone, cochlea, retina, endothelial cells, endometrium, pancreas and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the
25 disorder.

 The tissue distribution and homology to proline-rich proteins indicates that the protein is a extracellular matrix protein or an ingredient of bodily fluid. Polypeptides and polynucleotides corresponding to Gene NO: 19 are useful for cancer metastasis intervention, tissue culture additive, bone modeling, wound healing and tissue repair.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 20

 Gene NO: 20 is expressed primarily in prostate cancer, leukocytes, meningima, adult liver, pancreas, brain, and to a lesser extent in lung.

 Therefore, polynucleotides or polypeptides of the invention are useful as
35 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, prostate cancers. Similarly, polypeptides and antibodies directed to these

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate and brain, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., prostate, leukocytes, meningioma, liver, brain, pancreas and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

Prostate cancer cell lines are known to be responsive to estrogen and androgen. The protein expression of Gene NO: 20 appears to be influenced by both estrogen and androgen levels. The prostate cancer tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 20 are useful in the intervention and detection of prostate hyperplasia and prostate cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 21

The translation product of Gene NO: 21 is identical to the human wnt-7a gene. Wnt-7a is a secreted signaling molecule, thought to be important in signaling and the regulation of cell fate and pattern formation during embryogenesis. Specifically, knock out studies in mice have demonstrated that wnt7a plays a critical role in the development of the dorsal-ventral patterning in the developing limb, and to a lesser extent plays a role in the development of anterior-posterior patterning. Overexpression of wnt7a can induce transformation of cultured mammary cells, suggesting that it is an oncogene.

Expression of Gene NO: 21 has only been observed in testes.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, testicular cancer; abnormal limb development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the testes or developing embryo. For a number of disorders of the above tissues or cells, particularly of the developing embryo, expression of this gene at significantly higher or lower levels may routinely be detected in the developing embryo or amniotic fluid taken from a pregnant individual and compared relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Also, expression of this gene at significantly higher or lower levels may routinely be detected in the testes of patient suffering from testicular cancer and compared relative to the

standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to mouse wnt7a indicates that polypeptides and polynucleotides corresponding to Gene NO: 21 are useful to restore abnormal limb development in an affected individual. Furthermore, its oncogenic potential and tissue distribution indicates that it could serve as a diagnostic for testicular cancer.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 154 as residues: Gly-22 to Arg-28.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 22

Gene NO: 22 is expressed primarily in fetal liver/spleen, breast, testes and placenta and to a lesser extent in brain, and a series of cancer tissues.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, immune disorders, brain diseases, male infertility, and disposition to pregnant miscarriages. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, hematopoietic system, and sexual organs, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., liver, spleen, testes, placenta, and brain, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of this gene indicates that polypeptides and polynucleotides corresponding to Gene NO: 22 are useful as a marker for non-differentiated, dividing cells and hence could serve as an oncogenic marker. Its high expression in fetal liver, suggests an involvement in hematopoiesis and/or the immune system. Hence it is useful as a factor to enhance an individuals immune system, e.g., in individuals with immune disorders. It is also thought to affect the survival, proliferation, and differentiation of a number of hematopoietic cell lineages, including hematopoietic stem cells. Its disruption, e.g., mutation or altered expression, may also be a marker of immune disorder. Its expression in the testes, suggests it may be important in controlling male fertility. Expression of this gene in breast further reflects a

role in immune function and immune surveillance (breast lymph node). This gene is believed to be useful as a marker for breast cancer.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 155 as residues: Gln-57 to Lys-70 and Ala-91 to Pro-100.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 23

Gene NO: 23 is expressed primarily in bone marrow and brain (whole and fetal).

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, neurological, immune and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous and hematopoietic systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., bone marrow, brain, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 23 are useful in the diagnosis and treatment of disorders related to the central nervous system (e.g. neuro-degenerative conditions, trauma, and behavior abnormalities) and hematopoiesis. In addition, the expression in fetal brain indicates a role for this gene product in diagnosis of predisposition to developmental defects of the brain.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 156 as residues: Thr-23 to Tyr-29.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 24

Gene NO: 24 is expressed primarily in smooth muscle, placenta, prostate, and osteoblasts.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cardiovascular pathologies. Similarly, polypeptides and antibodies

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directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular, reproductive and skeletal systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., placenta, smooth muscle, prostate, and osteoblasts, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 24 are useful for detection and treatment of neoplasias and developmental abnormalities associated with these tissues.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 157 as residues: Asn-21 to Thr-26.

FEATURES OF PROTEIN ENCODED BY GENE NO: 25

The translation product of Gene NO: 25 shares sequence homology with Pregnancy Associated Mouse Protein (PAMP)-1. (See, FEBS Lett 1993 May 17;322(3):219-222). Based on the sequence similarity the translation product of this gene is expected to share certain biological activities with PAMP-1.

Gene NO: 25 is expressed primarily in 12-week-old human embryos and prostate.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, prostate disorders (cancer). Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., embryonic tissue, and prostate, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 25 are useful for the diagnosis and treatment of prostate disorders (such as cancer) and developmental abnormalities and fetal deficiencies. The homology to PAMP-1 indicates that this gene and gene product are useful in detecting pregnancy.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 158 as residues: Pro-23 to Glu-28 and Ser-44 to Gly-55.

FEATURES OF PROTEIN ENCODED BY GENE NO: 26

Gene NO: 26 is expressed primarily in testes and to a lesser extent in epididymis.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, reproductive and endocrine disorders, as well as testicular cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive and endocrine systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., testes, and epididymis, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 26 are useful for the treatment and diagnosis of conditions concerning proper testicular function (e.g., endocrine function, sperm maturation), as well as cancer. Therefore, this gene product is useful in the treatment of male infertility and/or impotence. This gene product is also useful in assays designed to identify binding agents as such agents (antagonists) are useful as male contraceptive agents. Similarly, the protein is believed to be useful in the treatment and/or diagnosis of testicular cancer. The testes are also a site of active gene expression of transcripts that may be expressed, particularly at low levels, in other tissues of the body. Therefore, this gene product may be expressed in other specific tissues or organs where it may play related functional roles in other processes, such as hematopoiesis, inflammation, bone formation, and kidney function, to name a few possible target indications.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 15 as residues: Pro-24 to Gly-33 and Arg-70 to Gly-76.

FEATURES OF PROTEIN ENCODED BY GENE NO: 27

5 The translation product of Gene NO: 27 shares sequence homology with salivary protein precursors which are thought to be important in immune response and production of secreted proteins.

Gene NO: 27 is expressed primarily in salivary gland tissue.

10 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, immune disorders, diseases of the salivary gland. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a
15 number of disorders of the above tissues or cells, particularly of the immune system, digestive system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., salivary gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder,
20 relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

 The tissue distribution and homology to salivary secreted protein indicates that polypeptides and polynucleotides corresponding to Gene NO: 27 are useful for treatment of immune disorders and diagnostic uses related to secretion of protein in
25 disease states. For example, the gene product can be used as an anti-microbial agent, an ingredient for oral or dental hygiene, treatment of xerostomia, sialorrhea, intervention for inflammation including parotitis, and an indication for tumors in the salivary gland (adenomas, carcinomas).

 Preferred epitopes include those comprising a sequence shown in SEQ ID NO:
30 160 as residues: Asp-21 to Gly-28, Asp-30 to Glu-43, Glu-49 to Glu-62, and Thr-75 to Pro-83.

FEATURES OF PROTEIN ENCODED BY GENE NO: 28

35 Gene NO: 28 is expressed primarily in human fetal heart tissue and to a lesser extent in olfactory tissue.

 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions which include, but are not limited to, immune, olfactory and cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, olfactory and vascular systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., olfactory tissue, and heart, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 28 are useful for diagnosis and treatment of immune, olfactory and vascular disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 161 as residues: Cys-33 to Gly-44, Arg-71 to Arg-78, Ser-130 to Gly-142, Lys-150 to Gly-157, and Thr-159 to Asp-177.

FEATURES OF PROTEIN ENCODED BY GENE NO: 29

Gene NO: 29 is expressed primarily in brain and to a lesser degree in activated macrophages, endothelial and smooth muscle cells, and some bone cancers.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of brain and endothelial present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, neurodegeneration, inflammation and other immune disorders, fibrotic conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification brain, smooth muscle, and endothelium. For a number of disorders of the above tissues or cells, particularly of the brain and endothelium, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues or cell types (e.g., brain, endothelial cells, macrophages, smooth muscle, and bone, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

Tissue distribution suggests polypeptides and polynucleotides corresponding to Gene NO: 29 are useful in study and treatment of neurodegenerative and immune disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 162 as residues: Asn-18 to Glu-20, Ser-33 to Gln-48, Cys-55 to Ser-56, Pro-67 to Cys-69.

FEATURES OF PROTEIN ENCODED BY GENE NO: 30

Gene NO: 30 is expressed primarily in early stage human brain and to a lesser extent in cord blood, heart, and some tumors.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of developing CNS tissue present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cardiovascular and neurodegenerative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., brain and heart, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that that polypeptides and polynucleotides corresponding to Gene NO: 30 are useful for the treatment of cancer and of neurodegenerative and cognitive disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 31

Gene NO: 31 is expressed primarily in brain and thymus and to a lesser extent in several other organs and tissues including the hematopoietic system, liver skin and bone

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, CNS disorders, hematopoietic system disorders, disorders of the endocrine system, bone, and skin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential

identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly CNS disorders, hematopoietic system disorders, disorders of the endocrine system, bone, and skin, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g.,

5 hematopoietic cells, brain, thymus, liver, bone, and epidermis, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

10 The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 31 are useful for treatment and diagnosis of CNS disorders, hematopoietic system disorders, disorders of the endocrine system, and of bone and skin.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 15 164 as residues: Thr-35 to Arg-40, Pro-55 to His-75, Pro-93 to Ala-98, Ala-111 to Pro-119, and Pro-132 to Glu-138.

FEATURES OF PROTEIN ENCODED BY GENE NO: 32

Gene NO: 32 is expressed primarily in organs and tissue of the nervous system 20 and to a lesser extent in various developing tissues and organs.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, disorders of the central nervous system and disorders of developing and 25 growing tissues and organs. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly disorders of the CNS, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., tissue of the nervous 30 system and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

35 The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 32 are useful for diagnosis and treatment of disorders of the central nervous system, general neurological diseases and neoplasias.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 165 as residues: Ser-33 to Lys-41 and Glu-86 to Glu-91.

FEATURES OF PROTEIN ENCODED BY GENE NO: 33

- 5 Residues 141-156 in the translation product for Gene NO: 33 as shown in the sequence listing matches phosphopantetheine binding site motifs. Phosphopantetheine (or pantetheine 4' phosphate) is the prosthetic group of acyl carrier proteins (ACP) in some multienzyme complexes where it serves as a 'swinging arm' for the attachment of activated fatty acid and amino-acid groups. Phosphopantetheine is attached to a serine
- 10 residue in these proteins. ACP proteins or domains have been found in various enzyme systems which are listed below. Fatty acid synthetase (FAS), which catalyzes the formation of long-chain fatty acids from acetyl-CoA, malonyl-CoA and NADPH. Bacterial and plant chloroplast FAS are composed of eight separate subunits which correspond to the different enzymatic activities; ACP is one of these polypeptides.
- 15 Fungal FAS consists of two multifunctional proteins, FAS1 and FAS2; the ACP domain is located in the N-terminal section of FAS2. Vertebrate FAS consists of a single multifunctional enzyme; the ACP domain is located between the beta-ketoacyl reductase domain and the C-terminal thioesterase domain. Based on the presence of a phosphopantetheine binding site in the translation product of this gene, it is believed to
- 20 share activities fatty acid synthetase polypeptides. Such activities may be assayed by methods known in the art.

This gene is expressed primarily in developing and rapidly growing tissues like placenta fetal heart and endometrial tumor and to a lesser extent in B and T cell lymphoma tissues

- 25 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer and disorders of developing tissues and organs. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing
- 30 immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic tissues and developing organs and tissues, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., embryonic tissue, endometrium, B-cells, and T-cells, and cancerous and wounded
- 35 tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to

the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 33 are useful for treatment and diagnosis of cancer in the hematopoietic system developing organs and tissues. It may also be useful for induction of cell growth in disorders of the hematopoietic system and other tissue and organs. The homology to fatty acid synthetases indicates that this gene product is useful in the diagnosis and treatment of lipid metabolism disorders such as hyperlipidemia.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 166 as residues: Arg-27 to Glu-34.

FEATURES OF PROTEIN ENCODED BY GENE NO: 34

Gene NO: 34 is expressed primarily in breast and testes tissues and to a lesser extent in hematopoietic tissues including tonsils, T cells and monocytes.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, diseases of the reproductive organs and systems, including cancer, autoimmune diseases and inflammatory diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive organs and hematopoietic tissues, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., hematopoietic cells, T-cells and monocytes, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Nucleic acids comprising sequence of this gene are also useful as chromosome markers since this gene maps to Chr. 15, D15S118-D15S123.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 34 are useful for treatment of diseases of the reproductive organs and hematopoietic system including cancer, autoimmune diseases and inflammatory diseases.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 167 as residues: Phe-81 to Lys-86.

FEATURES OF PROTEIN ENCODED BY GENE NO: 35

The translation product of Gene NO: 35 shares sequence similarity with the mouse cytokine-inducible inhibitor of signaling. See, e.g., Nature 1997 Jun 26;387(6636):917-921. Cytokines are secreted proteins that regulate important cellular responses such as proliferation and differentiation. Key events in cytokine signal transduction are well defined: cytokines induce receptor aggregation, leading to activation of members of the JAK family of cytoplasmic tyrosine kinases. In turn, members of the STAT family of transcription factors are phosphorylated, dimerize and increase the transcription of genes with STAT recognition sites in their promoters. Less is known of how cytokine signal transduction is switched off. Expression of the mouse SOCS-1 protein inhibited both interleukin-6- induced receptor phosphorylation and STAT activation. We have also cloned two relatives of SOCS-1, named SOCS-2 and SOCS-3, which together with the previously described CIS form a new family of proteins. Transcription of all four SOCS genes is increased rapidly in response to interleukin-6, in vitro and in vivo, suggesting they may act in a classic negative feedback loop to regulate cytokine signal transduction. The translation product of this gene is believed to have similar biological activities as this family of mouse genes. The biological activity of the translation product of this gene may be assayed by methods shown in Nature 1997 Jun 26;387(6636): 917-921, which is incorporated herein by reference in its entirety.

Gene NO: 35 is expressed primarily in tissues of hematopoietic origin including activated monocytes, neutrophils, activated T-cells and to a lesser extent in breast, adipose tissue and dendritic cells.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, diseases of the hematopoietic system including cancer autoimmune diseases and inflammatory diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic system expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., hematopoietic cells and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene

expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to cytokine inducible inhibitor of signaling indicates that polypeptides and polynucleotides corresponding to Gene NO: 35 are
5 useful for diagnosis and treatment of diseases of the hematopoietic system including autoimmune diseases, inflammatory diseases, infectious diseases and neoplasia. For example, administration of, or upregulation of this gene could be used to decrease the response of immune-system to lymphokines and cytokines.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO:
10 168 as residues: Arg-23 to His-30, Ala-35 to Gly-42.

FEATURES OF PROTEIN ENCODED BY GENE NO: 36

Gene NO: 36 is expressed primarily in infant brain and to a lesser extent in osteoclastoma, placenta, and a wide variety of other tissues.

15 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential
20 identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., osteoclastoma, placenta, and tissue of the central nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial
25 fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 36 are useful for diagnosis and treatment of neurologic
30 disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 169 as residues: Gln-31 to Ser-37, Ile-49 to Gly-54, Tyr-57 to Asp-67, Gln-141 to Pro-151, and Val-207 to Thr-219.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 37

Gene NO: 37 is expressed primarily in osteoclastoma stromal cells, dendritic cells, liver, and placenta.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer, wound, pathological conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues or cell types (e.g., stromal cells, dendritic cells, liver, and placenta and, cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 37 are useful for fundamental role in basic growth and development of human.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 170 as residues: Leu-32 to Thr-37 and Arg-48 to Pro-55.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 38

The translation product of Gene NO: 38 shares sequence homology with a yeast protein, Lpe10p, which may be involved in mRNA processing. (See Accession Nos. 2104457 and 1079682.) It is likely that an upstream signal sequence exists, other than the predicted sequence described in Table 1. Preferred polypeptide fragments comprise the open reading frame upstream from the predicted signal sequence, as well as polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in skin, and to a lesser extent in embryonic tissues, and fetal liver.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, defects of the skin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skin, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., epidermis, liver, and embryonic tissues, and cancerous and wounded tissues) or bodily fluids (e.g., serum,

plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

- 5 The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 38 are useful for diagnosis and treatment of defects of the skin.

FEATURES OF PROTEIN ENCODED BY GENE NO: 39

- 10 Gene NO: 39 is expressed primarily in Amygdala, activated monocytes, testis, and fetal liver. Moreover, this gene is mapped to chromosome 4. Thus, polynucleotides of the present invention can be used in linkage analysis as markers for chromosome 4.

- Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, defects of the brain, immune system and testis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, immune system and testis, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., Amygdala, monocytes, testes, and liver and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

- The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 39 are useful for detecting defects of the brain, immune system and testis because of its abundance in these tissues.

30

FEATURES OF PROTEIN ENCODED BY GENE NO: 40

- The translation product of Gene NO: 40 shares sequence homology with lymphoma 3-encoded protein (bcl-3) which is thought to contribute to leukemogenesis when abnormally expressed.
- 35 This gene is expressed primarily in Human Neutrophils, and to a lesser extent in Human Osteoclastoma Stromal Cells (unamplified), Hepatocellular Tumor, and Human Neutrophils, (Activated).

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, chronic lymphocytic leukemia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., neutrophils, osteoclastoma, and kidney, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to lymphoma 3-encoded protein (bcl-3) indicates that polypeptides and polynucleotides corresponding to Gene NO: 40 are useful for treatment of lymphoma and related cancers.

FEATURES OF PROTEIN ENCODED BY GENE NO: 41

Gene NO: 41 is expressed primarily in ovary tumor, and to a lesser extent in endometrial stromal cells and fetal brain.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, ovarian or endometrial cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the female reproductive system and the developing central nervous system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., ovary, endometrium and brain, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 41 are useful for development of factors involved in ovarian or endometrial and general reproductive organ disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 174 as residues: Glu-22 to Trp-31, Asn-84 to Asp-90, and Ser-144 to Asp-151.

FEATURES OF PROTEIN ENCODED BY GENE NO: 42

The translation product of Gene 42 has sequence identity with a gene designated PTHrP(B). The PTHrP(B) polypeptide inhibits parathyroid hormone related peptide (PTHrP) activity.

This gene is expressed primarily in adult testis, and to a lesser extent in pituitary.

Therefore polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of male reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., testes, and pituitary, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Furthermore, based in part on sequence identity with PTHrP(B), nucleic acids and polypeptides of the present invention may be used to diagnose or treat such conditions as hypercalcemia, osteoporosis, and disorders related to calcium metabolism.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 42 are useful for treatment of male reproductive disorders, hypercalcemia, osteoporosis, and other disorders related to calcium metabolism.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 175 as residues: Tyr-81 to Met-86, Gly-103 to Ser-108, Glu-127 to Pro-128, Pro-175 to Ser-180, Glu-196 to Lys-203, Pro-235 to Ser-241, and Ala-249 to Ser-264.

FEATURES OF PROTEIN ENCODED BY GENE NO: 43

The translation product of Gene NO: 43 shares sequence homology with brevican, which is thought to be important as a proteoglycan core protein of the

aggrecan/versican family. The translation product of this gene may also contain a hyaluronan (HA)-binding region domain in frame with, but downstream of, the predicted open reading frame (Barta, et al., Biochem. J. 292:947-949 (1993)). The HA-binding domain, also termed the link domain, is found in proteins of vertebrates that are involved in the assembly of extracellular matrix, cell adhesion, and migration. It is about 100 amino acids in length. The structure has been shown to consist of two alpha helices and two antiparallel beta sheets arranged around a large hydrophobic core similar to that of C-type lectin. This domain typically contains four conserved cysteines involved in two disulfide bonds.

10 This gene is expressed primarily in early stage human brain and to a lesser extent in frontal cortex and epileptic tissues.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of disorders associated with, or observed during, neuronal development. Similarly, polypeptides and antibodies directed to these polypeptides are useful as immunological probes for differential identification of neuronal and associated tissues and cell types. For a number of disorders of the above tissues or cells, particularly for those of the nervous system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., brain and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

25 The tissue distribution and homology to brevican indicates that polypeptides and polynucleotides corresponding to Gene NO: 43 are useful for neuronal regulation and signaling. The uses include directing or inhibiting axonal growth for the treatment of neuro-fibromatosis and in detection of glioses.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 176 as residues: Asp-28 to Arg-33 and Arg-126 to Arg-131.

FEATURES OF PROTEIN ENCODED BY GENE NO: 44

Gene NO: 44 is the human homolog of Notch-2 (Accession No. 477495) and mouse EGF repeat transmembrane protein (Accession No. 1336628), both genes are important in differentiation and development of an organism. The EGF repeat transmembrane protein is regulated by insulin like growth factor Type I receptor. These proteins are involved in cell-cell signaling and cell fate determination. Based on

homology, it is likely that this gene products also involved in cell differentiation and development. Although the predicted signal sequence is indicated in Table 1, it is likely that a second signal sequence is located further upstream. Moreover, further translated coding regions are likely found downstream from the disclosed sequence, which can easily be obtained using standard molecular biology techniques. A frameshift occurs somewhere around nucleotide 714, causing a frame shift in amino acid sequence from frame +2 to frame +3. However, using the homology of Notch-2 and EGF repeat transmembrane protein, the complete open reading frame can be elucidated. Preferred polynucleotide fragments comprise nucleotides 146-715, 281-715, and 714-965. Other preferred polypeptide fragments comprise the following EGF-like motifs: CRCASGFTGEDC (SEQ ID NO:260), CTCQVGFTGKEC (SEQ ID NO:261), CLNLPGSYQCQC (SEQ ID NO:262), CKCLTGFTGQKC (SEQ ID NO:263), and CQCLQGFTGQYC (SEQ ID NO:264).

Gene NO: 44 is expressed primarily in placenta and to a lesser extent in stromal and immune cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, hemophilia and other blood disorders, central nervous system disorders, muscle disorders, and any other disorder resulting from abnormal development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, hematopoietic and vascular systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., placenta, stromal and immune cells and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Notch-2 indicates that polypeptides and polynucleotides corresponding to Gene NO: 44 are useful for diagnosing and treating disorders relating to abnormal regulation of cell fate, induction, and differentiation of cells (e.g., cancer), epidermal growth factors, axonal pathfinding, and hematopoiesis.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 177 as residues: Gln-27 to Tyr-32, His-45 to Glu-55, Tyr-61 to Gly-77, Glu-99 to Ser-106, Ser-125 to Cys-131, and Thr-138 to Trp-144.

FEATURES OF PROTEIN ENCODED BY GENE NO: 45

The translation product of this gene shares sequence homology with Laminin A which is thought to be important in the binding of epithelial cells to basement membrane and is associated with tumor invasion. Moreover, the translated protein is homologous to the *Drosophila* LAMA gene (Accession No. 1314864), a gene expressed in the first optic ganglion of *Drosophila*. Thus, it is likely that the gene product from this gene is involved in the development of the eye. Nucleotide fragments comprising nucleotides 822-1223, 212-475, 510-731, and 1677-1754 are preferred. Also preferred are the polypeptide fragments encoded by these polynucleotide fragments. It is likely that a frame shift occurs somewhere between nucleotides 475 to 510, shifting the open reading frame from +2 to +3. However, the open reading frame can be clarified using known molecular biology techniques.

This gene is expressed primarily in human testes tumor and to a lesser extent in placenta and activated monocytes.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, invasive cancers or tumors of the epithelium, as well as disorders relating to eye development. Similarly, polypeptides and antibodies directed to these polypeptides are useful as immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of neoplastic conditions, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., testes, placenta, and monocytes and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Laminin A indicates that polypeptides and polynucleotides corresponding to Gene NO: 45 are useful for study and diagnosis of malignant or benign tumors, fibrotic disorders, and eye disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 178 as residues: Met-1 to Gly-8, Glu-32 to Ala-37, Met-113 to Asn-119, and Glu-139 to Gln-153.

FEATURES OF PROTEIN ENCODED BY GENE NO: 46

The translation product of Gene NO: 46 is novel and shares sequence homology with the product of the *Drosophila* tissue polarity gene frizzled. In vertebrates, it appears that there is a family of proteins that represent frizzled gene homologs. (See, 5 e.g., Accession Nos. 1946343 and AFO17989.) The *Drosophila* frizzled protein is thought to transmit polarity signals across the plasma membrane of epidermal cells. The structure of frizzled proteins suggest that they may function as a G-protein-coupled receptor. The frizzled proteins are thought to represent receptors for Wnt gene products - secreted proteins that control tissue differentiation and the development of embryonic 10 and adult structures. Inappropriate expression of Wnts has also been demonstrated to contribute to tumor formation. Moreover, mammalian secreted frizzled related proteins are thought to regulate apoptosis. (See Accession No. AFO17989.) The human homolog has also been recently cloned by other groups. (See Accession No. H2415415.) Thus, the protein encoded by this gene plays a role in mediating tissue 15 differentiation, proliferation, tumorigenesis and apoptosis. Preferred polypeptide fragments lack the signal sequence as described in Table 1, as well as N-terminal and C-terminal deletions. Preferred polynucleotide fragments encode these polypeptide fragments.

Gene NO: 46 is expressed primarily in fetal tissues - particularly fetal lung - and 20 adult cancers, most notably pancreas tumor and Hodgkin's lymphoma. Together, this distribution is consistent with expression in tissues undergoing active proliferation. The gene is also expressed to a lesser extent in other organs, including stomach, prostate, and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as 25 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer (particularly pancreatic cancer and/or Hodgkin's lymphoma), as well as other forms of aberrant cell proliferation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for 30 differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and hyperproliferative disorders, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., fetal tissue, pancreas, and tissue of the immune system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, 35 urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to frizzled indicates that polypeptides and polynucleotides corresponding to Gene NO: 46 are useful for influencing cell proliferation, differentiation, and apoptosis. The full-length protein or a truncated domain could potentially bind to and regulate the function of specific factors, such as Wnt proteins or other apoptotic genes, and thereby inhibit uncontrolled cellular proliferation. Expression of this protein within a cancer - such as via gene therapy or systemic administration - could effect a switch from proliferation to differentiation, thereby arresting the progression of the cancer.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 179 as residues: Pro-31 to Arg-37.

FEATURES OF PROTEIN ENCODED BY GENE NO: 47

The translation product of Gene NO: 47 shares sequence homology with members of the Rh/T2/S-glycoprotein family of ribonuclease-encoding genes. These ribonuclease proteins are found predominantly in fungi, plants, and bacteria and have been implicated in a number of functions, including phosphate-starvation response, self-incompatibility, and responses to wounding. A second group has recently cloned this same gene, calling it a ribonuclease 6 precursor. (See Accession No. 2209029.) This group also mapped the gene to chromosome 6, thus, the polynucleotides of the present invention can be used in linkage analysis as a marker for chromosome 6.

Gene NO: 47 is expressed primarily in hematopoietic cells and tissues, including macrophages, eosinophils, CD34 positive cells, T-cells, and spleen. It is also expressed to a lesser extent in brain and spinal cord.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, tumors of a hematopoietic origin, graft rejection, wounding, inflammation, and allergy. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., hematopoietic cells, and tissues and cells of the immune system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder,

relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the Rh/T2/S-glycoprotein family of ribonuclease-encoding genes indicates that polypeptides and polynucleotides
5 corresponding to Gene NO: 47 are useful as a cytotoxin that could be directed against specific cell types (e.g. cancer cells; HIV- infected cells), and that would be well tolerated by the human immune system.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 180 as residues: Ala-24 to Asp-30, Ile-51 to Tyr-61, Pro-69 to Ser-78, Pro-105 to Phe-
10 110, Asn-129 to Phe-135, Pro-187 to Glu-192, Lys-205 to Gln-224, and Pro-250 to His-256.

FEATURES OF PROTEIN ENCODED BY GENE NO: 48

The translation product of Gene NO: 48 shares sequence homology with
15 dolichyl-phosphate glucosyltransferase, a transmembrane-bound enzyme of the endoplasmic reticulum which is thought to be important in N-linked glycosylation, by catalyzing the transfer of glucose from UDP-glucose to dolichyl phosphate. (See Accession No. 535141.) Based on homology, it is likely that this gene product also play a role similar in humans. Preferred polynucleotide fragments comprise nucleotides
20 132-959. Also preferred are the polypeptide fragments encoded by this nucleotide fragment.

Gene NO: 48 is expressed primarily in endothelial cells and to a lesser extent in hematopoietic cells and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as
25 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, defects in proper N-linked glycosylation of proteins, such as Wiskott-Aldrich syndrome; tumors of an endothelial cell origin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes
30 for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular and hematopoietic systems, as well as brain, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., endothelial cells, hematopoietic cells, and brain, and cancerous and wounded tissues) or bodily fluids
35 (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene

expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to dolichyl-phosphate glucosyltransferase indicates that polypeptides and polynucleotides corresponding to Gene NO: 48 are
5 useful in diagnosing and treating defects in N-linked glycosylation pathways that contribute to disease conditions and/or pathologies.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 181 as residues: Lys-50 to Thr-55, Ser-73 to Arg-79, Glu-92 to Pro-99, Asp-110 to Ser-117, Gln-125 to Lys-131, Gly-179 to Asn-188, Ile-231 to Cys-236, and Glu-318
10 to Asn-324.

FEATURES OF PROTEIN ENCODED BY GENE NO: 49

Gene NO: 49 is expressed primarily in brain, most notably in the hypothalamus and amygdala. This gene is also mapped to chromosome X, and therefore, can be used
15 in linkage analysis as a marker for chromosome X.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, tumors of a brain origin; neurodegenerative disorders, and sex-linked
20 disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., brain and cancerous and wounded tissues) or bodily
25 fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides
30 corresponding to Gene NO: 49 are useful for the diagnosis of tumors of a brain origin, and the treatment of neurodegenerative disorders, such as Parkinson's disease, and sex-linked disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 50

35 The translation product Gene NO: 50 shares sequence homology with canine phospholemman, a major plasma membrane substrate for cAMP-dependent protein kinases A and C. (See Accession No. M63934; see also Accession No. A40533.) In

fact, a group also recently cloned the human phospholemman gene, and mapped this gene to chromosome 19. (See Accession No.1916010.) Phospholemman is a type I integral membrane protein that gets phosphorylated in response to specific extracellular stimuli such as insulin and adrenalin. Phospholemman forms ion channels in the cell membrane and appears to regulate taurine transport, suggesting an involvement in cell volume regulation. It has been proposed that phospholemman is a member of a superfamily of membrane proteins, characterized by single transmembrane domains, which function in transmembrane ion flux. They are capable of linking signal transduction to the regulation of such cellular processes as the control of cell volume.

Gene No 50 is expressed primarily in fetal liver and to a lesser extent in adult brain and kidney, as well as other organs.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, insulin and/or adrenalin defects; diabetes; aberrant ion channel signaling; defective taurine transport; and defects in cell volume regulation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and/or immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., liver, brain, and kidney, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to phospholemman indicates that polypeptides and polynucleotides corresponding to Gene NO: 50 are useful for treatment of disorders involving the transport of ions and small molecules, in particular taurine. It could also be beneficial for control of pathologies or diseases wherein aberrancies in the control of cell volume are a distinguishing feature, due to the predicted role for phospholemman in the normal control of cell volume. It also may play a role in disorders involving abnormal circulating levels of insulin and/or adrenalin - along with other active secreted molecules - as revealed by its phosphorylation upon stimulation with insulin or adrenalin.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 183 as residues: Ala-20 to Gln-34, Arg-58 to Thr-79, and Leu-87 to Arg-92.

FEATURES OF PROTEIN ENCODED BY GENE NO: 52

Gene NO: 52 is expressed primarily in metastatic melanoma and to a lesser extent in infant brain.

5 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer and cancer metastasis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for
10 differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., epidermis, and brain, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a
15 disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

 The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 52 are useful for diagnosis and treatment of melanoma.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 53

 The translation product of Gene NO: 53 shares sequence homology with mucin which is thought to be important cell surface molecule. It also exhibits sequence identity with a calcium channel blocker of Agelenopsis aperta. In particular, with those calcium channel blockers which affect neuronal and muscle cells.

25 Gene NO: 53 is expressed primarily in prostate, endothelial cells, smooth muscle and fetal tissues and to a lesser extent in T cells and placenta.

 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are
30 not limited to, prostate cancer, immune disorders, angina, hypertension, cardiomyopathies, supraventricular arrhythmia, oesophageal achalasia, premature labour, and Raynaud's disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,
35 particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues or cell types (e.g., prostate, and tissue and cells of the immune system, and cancerous and wounded tissues) or

bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

- 5 The tissue distribution and homology to mucin indicates that polypeptides and polynucleotides corresponding to Gene NO: 53 are useful as a surface antigen for diagnosis of diseases such as prostate cancer and as tumor vaccine.

FEATURES OF PROTEIN ENCODED BY GENE NO: 54

- 10 Gene NO: 54 encodes a polypeptide which exhibits sequence identity with the rab receptor and VAMP-2 receptor proteins. (Martincic, et al., J. Biol. Chem. 272 (1997).)

Gene NO: 54 is expressed primarily in placenta, fetal liver, osteoclastoma and smooth muscle and to a lesser extent in T cell, fetal lung and colon cancer.

- 15 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancers, osteoporosis and immuno-related diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing
- 20 immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, hematopoiesis system and bone system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., placenta, liver, osteoclastoma, smooth muscle, T-cells, and lung, and colon, and
- 25 cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

- 30 The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 54 are useful for treating cancer, osteoporosis and immuno-disorders.

- Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 187 as residues: Pro-16 to Phe-21, Pro-24 to Arg-35, Arg-92 to Pro-98, Asn-143 to
- 35 Lys-151, and Leu-169 to Ile-176.

FEATURES OF PROTEIN ENCODED BY GENE NO: 55

Gene NO: 55 encodes a protein having sequence identity to the rat galanin receptor GALR2.

Gene NO: 55 is expressed primarily in ovarian cancer.

5 Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of ovarian cancer. Similarly, polypeptides and antibodies directed to those polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders
10 of the above tissues or cells, particularly of the immune system and reproductive system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., ovary, and tissues and cells of the immune system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell sample
15 taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. GALR2 antagonists can be used to treat obesity, bulimia, or Alzheimer's disease, while GALR2 agonists can be used to treat anorexia or pain, or to decrease nociception (claimed). Agonists and antagonists can also be used to
20 treat numerous other disorders, including cognitive disorders, sensory disorders, motion sickness, convulsion/epilepsy, hypertension, diabetes, glaucoma, reproductive disorders, gastric and intestinal ulcers, inflammation, immune disorders, and anxiety.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 55 are useful for diagnosis and treatment of ovarian cancer.
25

FEATURES OF PROTEIN ENCODED BY GENE NO: 56

As indicated in Table 1, the predicted signal sequence of Gene NO: 56 relates to an open reading frame that is homologous to the mouse major histocompatibility locus class III. (See Accession No. 2564953.) Any frame shift mutations that alter the correct
30 open reading frame can easily be clarified using known molecular biology techniques. Moreover, in the opposite orientation, a second translated product is disclosed. This second translation product of this contig is identical in sequence to intracellular protein lysophosphatidic acid acyltransferase. The nucleotide and amino acid sequences of this translated product have since been published by Stamps and colleagues (Biochem. J.
35 326 (Pt 2), 455-461 (1997)), West and coworkers (DNA Cell Biol. 6, 691-701 (1997)), Rowan (GenBank Accession No. U89336), and Soyombo and Hofmann (GenBank Accession No. AF020544). This gene is thought to enhance cytokine

signaling response in cells. It is likely that a signal peptide is located upstream from this translated product. Preferred polypeptide fragments comprise the amino acid sequence: GLACWLAGVIFIDRKRTGDAISVMSEVAQTLLTQDVXVWVFPEGTRNHNGSML PFKRGAFHLAVQAQVPIVPIVMSSYQDFYCKKERRFTSGQCQVRVLPVPTEGL
 5 TPDVPALADRVRHSMHLHCF(SEQ ID NO: 265);
 PSAKYFFKMAFYNGWILFLAVLAIPVCAVRGRNVENMKILRLMLLHIKYLYGI
 RVEVRGAHHFPPSQPYVVVSNHQSSDLLGMMEVLPGRVCVPIAKR (SEQ ID
 NO:266); TVFREISTD (SEQ ID NO:267); or LWAGSAGWPAG (SEQ ID NO: 268).
 Also provided are polynucleotide fragments encoding these polypeptide fragments.

10 Gene NO: 56 is expressed primarily in infant adrenal gland, hypothalamus, 7
 week old embryonic tissue, fetal lung, osteoclastoma stromal cells, and to a lesser
 extent in a large number of additional tissues.

Therefore, polynucleotides or polypeptides of the invention are useful as
 reagents for differential identification of the tissue(s) or cell type(s) present in a
 15 biological sample and for diagnosis of developmental disorders and osteoclastoma.
 Similarly, polypeptides and antibodies directed to these polypeptides are useful in
 providing immunological probes for differential identification of the tissue(s) or cell
 type(s) in which it is highly expressed. For a number of disorders of the above tissues
 or cells, particularly during development or of the nervous or bone systems, expression
 20 of this gene at significantly higher or lower levels may routinely be detected in certain
 tissues and cell types (e.g., adrenal, embryonic tissue, lung, and osteoclastoma stromal
 cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine,
 synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell
 sample taken from an individual having such a disorder, relative to the standard gene
 25 expression level, i.e., the expression level in healthy tissue or bodily fluid from an
 individual not having the disorder. Further, expression of this protein can be used to
 alter the fatty acid composition of a given cell or membrane type.

The tissue distribution indicates that polypeptides and polynucleotides
 corresponding to Gene NO: 56 are useful for diagnosis and treatment of osteoclastoma
 30 and other bone and non-bone-related cancers, as well as for the diagnosis and treatment
 of developmental disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO:
 189 as residues: Gly-29 to Gly-36 and Tyr-49 to Tyr-58.

35 **FEATURES OF PROTEIN ENCODED BY GENE NO: 57**

The translation product of Gene NO: 57 shares sequence homology with
 longevity-assurance protein-1. (See Accession No. g1123105.) Preferred

polynucleotide fragments comprise nucleotides 6-125 and 118-432, as well as the polypeptides encoded by these polynucleotides. It is likely that a second signal sequence exists downstream from the predicted signal sequence in Table 1. Moreover, a frame shift likely occurs between nucleotides 118-125, which can be elucidated using standard molecular biology techniques.

Gene NO: 57 is expressed primarily in fetal liver, kidney, brain, thymus, and bone marrow.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, immunological diseases and hyperproliferative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal liver, kidney, brain, thymus, and bone marrow expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., liver, kidney, brain, thymus, and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to longevity-assurance protein suggest that Gene NO: 57 encodes a protein useful in increasing life span and in replacement therapy for those suffering from immune system disorders or hyperproliferative disorders caused by underexpression or overexpression of this gene.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 190 as residues: Val-29 to Arg-46 and Gly-50 to Gly-56.

FEATURES OF PROTEIN ENCODED BY GENE NO: 58

Domains of the Gene NO: 58 product are homologous to porcine surfactant protein-A receptor. (See Accession No. B48516.) The bovine gene binds surfactant protein-A receptor, modulating the secretion of alveolar surfactant. Based on this homology, the gene product encoded by this gene will likely have activity similar to the porcine gene. Preferred polynucleotide fragments comprise nucleotides 887-1039, as well as the polypeptide fragments encoded by this nucleotide fragment.

Gene NO: 58 is expressed primarily in brain and to a lesser extent in endothelial cells.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, diseases of the central nervous system including dementia, stroke, neurological disorders, respiratory distress, and diseases affecting the endothelium including inflammatory diseases, restenosis, and vascular diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the placenta, liver, endothelial cells, prostate, thymus, and lung, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., brain, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology indicates that polypeptides and polynucleotides corresponding to Gene NO: 58 are useful for the diagnosis and /or treatment of diseases on the central nervous system, such as a factor that promote neuronal survival or protection, in the treatment of inflammatory disorders of the endothelium, or in disorders of the lung. In addition this protein may inhibit or promote angiogenesis and therefore is useful in the treatment of vascular disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 191 as residues: His-66 to Pro-80, Gly-139 to Ser-146 and Ser-262 to Pro-267.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 59

The translation product of Gene NO: 59 is homologous to the rat hypertension-induced protein which is thought to be important in hypertension, and found expressed mainly in kidneys. (See Accession No. B61209.) Thus, it is likely that this gene product is involved in hypertension in humans. Preferred polypeptide fragments comprise the short chain dehydrogenase/reductase motif SILGIISVPLSIGYCASKHALRGFFNGLR (SEQ ID NO:269), as well as polynucleotides encoding this polypeptide fragment. Also preferred are polynucleotide fragments of 337-639, as well as the polypeptide fragments encoded by this polynucleotide fragment.

35

Gene NO: 59 is expressed primarily in liver, spleen, lung, brain, and prostate.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cardiovascular, immunological, and renal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular, renal, and immune, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., liver, spleen, lung, brain, and prostate, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to hypertension-induced protein indicates that polypeptides and polynucleotides corresponding to Gene NO: 59 are useful for treating hypertension.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 192 as residues: Gln-40 to Glu-45, Glu-96 to Glu-102, Asn-256 to Thr-266, and Asp-308 to Asp-317.

FEATURES OF PROTEIN ENCODED BY GENE NO: 60

Gene NO: 60 is expressed primarily in activated T-cell and jurkat cell and to a lesser extent in apoptic T-cell and CD34+ cell. It is likely that alternative open reading frames provide the full length amino acid sequence, which can be verified using standard molecular biology techniques.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, T lymphocyte related diseases or hematopoiesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., T-cells and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard

gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 60 are useful for diagnosis or treatment of immune system disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 61

The translation product of Gene NO: 61, a vacuolar proton-ATPase, shares sequence homology with a *Caenorhabditis elegans* protein which is thought to be important in development. This protein may be a human secretory homologue that may also influence embryo development. Ludwig, J., also recently cloned this gene from chromaffin granules. (See, Accession No. 2584788.) Although Table 1 indicates the predicted signal peptide sequence, the translated product of this gene may in fact start with the upstream methionine, beginning with the amino acid sequence MAYHGLTV (SEQ ID NO:270). Thus, polypeptides comprising this upstream sequence, as well as N-terminus deletions, are also contemplated in the present invention.

Gene NO: 61 is expressed primarily in human placenta, liver, and Hodgkin's Lymphoma and to a lesser extent in bone marrow. Modest levels of expression were also observed in dendritic cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, hyperproliferative disorders, defects in embryonic development, and diseases or disorders caused by defects in chromaffin granules. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly cancer, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., placenta, liver, lymph tissue, and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to *Caenorhabditis elegans* indicates that polypeptides and polynucleotides corresponding to Gene NO: 61 are useful for diagnostic or therapeutic modalities for hyperproliferative disorders, embryonic development disorders, and chromaffin granules disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 62

The translation product of Gene NO: 62 shares sequence homology with the murine LAG3 gene which is thought to be important in the mediation of natural killer cell (NK cell) activity as previously determined by experiments in mice containing null mutations of LAG3. The similarity of this gene to the CD4 receptor may imply that the gene product may be a secreted, soluble receptor and immune mediator.

Gene NO: 62 is expressed primarily in human fetal heart, meningima, and to a lesser extent in tonsils. This gene also is expressed in the breast cancer cell line MDA 36.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, lymphomas, leukemias, breast cancer and any immune system dysfunction, including those dysfunctions which involve natural killer cell activities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system or breast cancer, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., heart, meningima, and tonsils and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the LAG3 gene (murine) indicates that the polynucleotides and polypeptides corresponding to Gene NO: 62 are useful for diagnostic and/or therapeutic modalities directed at abnormalities or disease states involving defective immune systems, preferably involving natural killer cell activity, as well as breast cancer.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 195 as residues: Pro-10 to Trp-17, Cys-58 to Pro-67, Thr-76 to Glu-85, and Arg-93 to Asn-101.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 63

The translation product of Gene NO: 63 shares sequence homology with a *Caenorhabditis elegans* alpha-collagen gene (Clg), which is thought to be important in

organism development, as well as other collagen genes. Thus, based on sequence homology, polypeptides of this gene are expected to have activity similar to collagen, including involvement in organ development.

Gene NO: 63 is expressed primarily in human B-Cell Lymphoma, and to a
5 lesser extent in human pituitary tissue. This gene has also demonstrated expression in keratinocytes.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are
10 not limited to, B-Cell Lymphoma, other lymphomas, leukemias, and other cancers, as well as disorders related to development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at
15 significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., tissue and/or cells of the immune system, and pituitary, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue
20 or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to *Caenorhabditis elegans* alpha-collagen gene indicates that polypeptides and polynucleotides corresponding to Gene NO: 63 are useful for development of diagnostic and/or therapeutic modalities directed at the detection and/or treatment of cancer, specifically B-Cell Lymphomas, leukemias, or
25 diseases related to development.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 196 as residues: Thr-22 to Arg-27 and Ser-29 to Thr-39.

FEATURES OF PROTEIN ENCODED BY GENE NO: 64

The translation product of Gene NO: 64 shares sequence homology with human
30 extracellular molecule olfactomedin, which is thought to be important in the maintenance, growth, or differentiation of chemosensory cilia on the apical dendrites of olfactory neurons. Based on this sequence homology, it is likely that polypeptides of this gene have activity similar to the olfactomedin, particularly the differentiation or
35 proliferation of neurons.

Gene NO: 64 is expressed primarily in fetal lung tissue.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, diseases of the lung as well as neural development, particularly of the lung. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the pulmonary system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., lungs and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the olfactomedin family indicates that polypeptides and polynucleotides corresponding to Gene NO: 64 are useful for the development of diagnostic and/or therapeutic modalities directed at detection and/or treatment of pulmonary disease states, e.g., cystic fibrosis.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 197 as residues: Gly-17 to Gln-23, Gln-45 to Arg-50, Arg-56 to Lys-61, Glu-70 to Leu-76, Asp-88 to Glu-93, Pro-117 to Met-131, Asp-161 to Glu-167, Arg-224 to Asn-237, Asp-302 to Trp-312, Pro-315 to Asn-320, and Thr-337 to Ser-341.

FEATURES OF PROTEIN ENCODED BY GENE NO: 65

The translation product of Gene NO: 65 shares sequence homology with *Saccharomyces cerevisiae* hypothetical protein YKL166 (Accession No. gi/687880) which is thought to be important in secretory and/or vesicular transport mechanisms. Based on this homology, it is likely that the gene product would have similar activity to YKL166, particularly secretory or transport mechanisms. Preferred polypeptide fragments of this gene include those fragments starting with the amino acid sequence ISAARV (SEQ ID NO:271). Other polypeptide fragments include the former fragment, which ends with the amino acid sequence PDVSEFMTRLF (SEQ ID NO:272). Further preferred fragments include those polypeptide fragments comprising the amino acid sequence FDPVRVDITSGKMRAR (SEQ ID NO:273). Also preferred are polypeptide fragments having exogenous signal sequences fused to the polypeptide.

Gene No 65 is expressed primarily in placenta, testis, osteoclastoma and to a lesser extent in adrenal gland.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer and/or diseases involving defects in protein secretion. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, cartilage and bone, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., placenta, testis, adrenal gland, and osteoclastoma, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the yeast YKL1GG protein indicates that polypeptides and polynucleotides corresponding to Gene NO: 65 are useful for the development of therapeutic and/or diagnostic modalities targeted at cancer or secretory anomalies, such as genetically caused secretory diseases.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 198 as residues: Ser-18 to Ser-29 and Lys-53 to Arg-74.

FEATURES OF PROTEIN ENCODED BY GENE NO: 66

The translation product of Gene NO: 66 shares sequence homology with the human papilloma virus (HPV) E5 ORF region which is thought to be important as a secreted growth factor. Although this is described as a viral gene product, it is believed to have several cellular secretory homologues. Therefore, based on the sequence similarity between the HPV E5 ORF and the translated product of this gene, this gene product is likely to have activity similar to HPV E5 ORF.

Gene NO: 66 is expressed primarily in activated T-Cells, monocytes, cerebellum and to a lesser extent in infant brain.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer and/or human papilloma virus infection. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of

this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., brain, lymph tissue, monocytes, and T-cells and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Moreover, polynucleotides of this gene have been mapped to chromosome 1. Therefore, polynucleotides of the present invention can be used in linkage analysis as a marker for chromosome 1.

The tissue distribution and homology to human papilloma virus E5 region indicates that polypeptides and polynucleotides corresponding to Gene NO: 66 are useful for development of diagnostic and/or therapeutic modalities directed at the diagnosis and/or treatment of cancer and/or human papilloma virus infection (HPV).

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 199 as residues: Asn-31 to Arg-36 and Leu-102 to Ser-112.

FEATURES OF PROTEIN ENCODED BY GENE NO: 67

The translation product of Gene NO: 67 shares sequence homology with the 8hs20 protein precursor [*Mus musculus*] which is thought to be important in B-Cell mu chain assembly. (See, Accession No. PID/d1002996; Shiraswa, T., EMBO. J. 12(5):1827-1834 (1993).) A polypeptide fragment starting at amino acid 53 is preferred, as well as 1-20 amino acid N-terminus and/or C-terminus deletions. Based on the sequence similarity between 8hs20 protein and the translation product of this gene, the two polypeptides are expected to share certain biological activities, particularly immunologic activities.

Gene NO: 67 is expressed primarily in human B-cells and to a lesser extent in Hodgkin's Lymphoma. It is also likely that the polypeptide will be expressed in B-cell specific cells, bone marrow, and spleen, as is observed with 8hs20.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, Hodgkin's Lymphoma, Common Variable Immunodeficiency, and/or other B-cell lymphomas. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., bone

marrow, spleen, lymph tissue, and B-cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to 8hs20 protein precursor [*Mus musculus*], indicates that polypeptides and polynucleotides corresponding to Gene NO: 67 are useful for therapeutic and/or diagnostic purposes, targeting Hodgkin's Lymphoma, B-cell lymphomas, Common Variable Immunodeficiency, or other immune disorders.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 200 as residues: Asp-51 to Trp-56, Arg-72 to Asp-85, and Gln-106 to Asp-112.

FEATURES OF PROTEIN ENCODED BY GENE NO: 68

Gene NO: 68 is expressed primarily in fetal liver/spleen, rhabdomyosarcoma, and to a lesser extent in 9 week-old early stage human embryo and bone marrow.

Therefore, polynucleotides or polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, rhabdomyosarcoma and other cancers, hematopoietic disorders, and immune dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues (e.g., embryonic tissue, striated muscle, liver, spleen, and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product of Gene NO: 68 is useful for diagnostic and/or therapeutic purposes directed to cancer, preferably rhabdomyosarcoma. Enhanced expression of this gene in fetal liver, spleen, and bone marrow indicates that this gene plays an active role in hematopoiesis. Polypeptides or polynucleotides of the present invention may therefore help modulate survival, proliferation, and/or differentiation of various hematopoietic lineages, including the hematopoietic stem cell. Thus, polynucleotides or polypeptides can be used treat

various hematopoietic disorders and influence the development and differentiation of blood cell lineages, including hematopoietic stem cell expansion. The polypeptide does contain a thioredoxin family active site at amino acids 64-82. Polypeptides comprising this thioredoxin active site are contemplated.

5

FEATURES OF PROTEIN ENCODED BY GENE NO: 69

Gene NO: 69 is expressed primarily in liver and kidney and to a lesser extent in macrophages, uterus, placenta, and testes.

Therefore, polynucleotides or polypeptides of the invention are useful as
10 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, renal disorders, neoplasms (e.g., soft tissue cancer, hepatocellular tumors), immune disorders, endocrine imbalances, and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in
15 providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic, urogenital, immune, and reproductive systems, expression of this gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., liver, kidney, uterus, placenta, testes, and macrophages and cancerous and
20 wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polypeptides and polynucleotides
25 corresponding to Gene NO: 69 are useful for diagnosis and treatment of disorders in the hepatic, urogenital, immune, and reproductive systems.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 202 as residues: Arg-41 to Ser-50, Glu-138 to Asn-148, Ser-155 to Arg-172, Pro-219 to Glu-228.

30

FEATURES OF PROTEIN ENCODED BY GENE NO: 70

Gene NO: 70 is expressed primarily in the immune system, including macrophages, T-cells, and dendritic cells and to a lesser extent in fetal tissue.

Therefore, polynucleotides or polypeptides of the invention are useful as
35 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, immune disorders, inflammatory diseases, lymph node disorders, fetal

development, and cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and hematopoietic systems expression of this gene at
5 significantly higher or lower levels may routinely be detected in certain tissues and certain cell types (e.g., macrophages, T-cells, dendritic cells, and fetal tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level
10 in healthy tissue or bodily fluid from an individual not having the disorder. There is some evidence that the polynucleotide is mapped to chromosome 19. Thus, the polynucleotide can be a marker for genetic analysis for chromosome 19.

The tissue distribution indicates that polypeptides and polynucleotides corresponding to Gene NO: 70 are useful for treatment, prophylaxis, and diagnosis of
15 immune and autoimmune diseases, such as lupus, transplant rejection, allergic reactions, arthritis, asthma, immunodeficiency diseases, leukemia, and AIDS. The polypeptides or polynucleotides of the present invention are also useful in the treatment, prophylaxis, and detection of thymus disorders, such as Graves Disease, lymphocytic thyroiditis, hyperthyroidism, and hypothyroidism. The expression observed
20 predominantly in hematopoietic cells also indicates that the polynucleotides or polypeptides are important in treating and/or detecting hematopoietic disorders, such as graft versus host reaction, graft versus host disease, transplant rejection, myelogenous leukemia, bone marrow fibrosis, and myeloproliferative disease. The polypeptides or polynucleotides are also useful to enhance or protect proliferation, differentiation, and
25 functional activation of hematopoietic progenitor cells (e.g., bone marrow cells), useful in treating cancer patients undergoing chemotherapy or patients undergoing bone marrow transplantation. The polypeptides or polynucleotides are also useful to increase the proliferation of peripheral blood leukocytes, which can be used in the combat of a range of hematopoietic disorders, including immunodeficiency diseases, leukemia, and
30 septicemia.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 203 as residues: Thr-21 to Ser-27, Pro-33 to Ser-38, and Arg-73 to Lys-84.

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of AA of F
1	HGCM20	97901 02/26/97 209047 05/15/97	pSport1	11	1739	25	1658	54	54	134	1	28	29	466
2	HLDBG33	97898 02/26/97 209044 05/15/97	pCMVSPORT 3.0	12	844	1	844	39	39	135	1	28	29	221
2	HLDBG33	97898 02/26/97 209044 05/15/97	pCMVSPORT 3.0	81	795	1	434	10	10	204	1	29	30	34
3	HTGEW86	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	13	776	134	676	173	173	136	1	35	36	155
4	HKCSR70	97900 02/26/97 209046 05/15/97	pBluescript	14	1376	727	1343	202	202	137	1	20	21	232
4	HKCSR70	97900 02/26/97 209046 05/15/97	pBluescript	82	1324	741	1309		861	205	1	31	32	42

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
4	HETBI87	209010 04/28/97 209085 05/29/97	Uni-ZAP XR	83	1494	1	1484	51	51	206	1	34	35	84
5	HTEAU17	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	15	502	1	502	143	143	138	1	33	34	60
6	HBMCY91	97897 02/26/97 209043 05/15/97	pBluescript	16	425	1	425	56	56	139	1	17	18	72
7	HSSGE07	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	17	1316	1	1298	45	45	140	1	26	27	376
7	HSSGE07	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	84	1285	1	1271	15	15	207	1	28	29	207
8	HMBX59	97897 02/26/97 209043 05/15/97	pBluescript	18	436	87	384	157	157	141	1	21	22	42

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
9	HNGIT22	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	19	503	1	503	23	23	142	1	19	20	40
10	HERAD57	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	20	358	1	358	147	147	143	1	31	32	69
11	HCENJ40	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	21	1926	573	1926	157	157	144	1	30	31	482
11	HCENJ40	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	85	394	1	394	166	166	208	1	20	21	23
11	HCENJ40	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	86	1925	573	1925	157	157	209	1	30	31	482
11	HCENJ40	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	87	1818	30	1298		1137	210	1			12

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of Sig OR F
12	HCSRA90	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	22	1224	64	557	80	80	145	1	30	31	225
13	HBJFC03	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	23	694	1	694	181	181	146	1	39	40	44
13	HBJFC03	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	88	539	1	539	215	215	211	1	18	19	19
14	HSNBL85	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	24	796	405	796	1	1	147	1	30	31	131
14	HSNBL85	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	89	855	300	855	513	513	212	1	37	38	54
15	HTEBY26	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	25	662	205	653	77	77	148	1	30	31	91

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
15	HTEBY26	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	90	628	198	625		275	213	1	31	32	34
16	HMA BH07	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	26	1105	40	1105	88	88	149	1	18	19	164
16	HMA BH07	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	91	1053	61	1009	79	79	214	1	22	23	229
17	HSKNY94	97899 02/26/97 209045 05/15/97	pBluescript	27	1017	1	1017	97	97	150	1	30	31	138
17	HSKNY94	97899 02/26/97 209045 05/15/97	pBluescript	93	2492	1	943	100	100	216	1	27	28	126
18	HMCDA67	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	28	391	1	391	169	169	151	1	29	30	57

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
19	HOSFF45	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	29	1139	6	1139	109	109	152	1	44	45	47
19	HOSFF45	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	94	3058	1795	2847	1868	1868	217	1	46	47	46
20	HMJAA51	97899 02/26/97 209045 05/15/97	pSport1	30	465	1	370	47	47	153	1	28	29	41
20	HMJAA51	97899 02/26/97 209045 05/15/97	pSport1	95	1099	664	1000	669	669	218	1	33	34	40
21	HTEBF05	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	31	702	1	702	403	403	154	1	24	25	71
22	HTEAL31	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	32	1142	1	518	49	49	155	1	47	48	105

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
22	HTEAL31	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	96	1580	23	422	32	32	219	1	47	48	104
23	HBMCT32	97899 02/26/97 209045 05/15/97	pBluescript	33	928	1	928	48	48	156	1	27	28	28
23	HBMCT32	97899 02/26/97 209045 05/15/97	pBluescript	97	678	72	593	89	89	220	1	27	28	28
24	HSKXE91	97899 02/26/97 209045 05/15/97	pBluescript	34	773	1	773	39	39	157	1	22	23	52
24	HSKXE91	97899 02/26/97 209045 05/15/97	pBluescript	98	1253	507	1253	507	507	221	1			16
25	HPWBTB39	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	35	453	1	453	40	40	158	1	25	26	74

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
26	HTLEV12	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	36	459	1	459	25	25	159	1	24	25	80
27	HSPAF93	97900 02/26/97 209046 05/15/97	pSport1	37	509	1	509	1	1	160	1	19	20	138
27	HSPAF93	97900 02/26/97 209046 05/15/97	pSport1	99	447	1	447	7	7	222	1	23	24	137
28	HHFGL62	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	38	598	1	598	1	1	161	1	21	22	177
28	HHFGL62	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	100	611	37	611	17	17	223	1	26	27	49
29	HCEIU14	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	39	454	1	454	1	1	162	1	21	22	71

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
29	HCEIU14	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	101	609	176	609	237	237	224	1			14
30	HEBDA39	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	40	425	1	376	223	223	163	1	18	19	66
31	HTHBA79	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	41	2471	141	2471	213	213	164	1	30	31	154
31	HTHBA79	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	102	1770	47	1721	119	119	225	1	31	32	154
31	HTHBA79	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	103	1832	96	1777	138	138	226	1			9
32	HAGBB70	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	42	2659	1172	2659	119	119	165	1	18	19	103

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
32	HAGBB70	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	104	2237	878	2237	1134	1134	227	1			19
33	HETDG84	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	43	1635	100	1580	299	299	166	1	20	21	80
34	HTEGA81	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	44	780	19	717	10	10	167	1	23	24	92
34	HKGAJ40	209236 09/04/97	pSport1	105	1822	1	1023	272	272	228	1	23	24	93
34	HKMLK44	209084 05/29/97	pBluescript	106	1712	1	1669	168	168	229	1	21	22	93
35	HTXAK60	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	45	2378	1337	2378	1437	1437	168	1	30	31	57
35	HTXAK60	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	107	1969	1068	1892	989	989	230	1	23	24	36

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
36	HMHBN40	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	46	1772	69	1772	129	129	169	1	30	31	231
36	HMHBN40	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	108	1734	65	1734	100	100	231	1	29	30	80
37	HFVGS85	97901 02/26/97 209047 05/15/97	pBluescript	47	1107	70	1107	83	83	170	1	30	31	71
38	HERAH81	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	48	805	167	764	167	167	171	1	23	24	64
39	HMSEU04	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	49	1408	131	1258	364	364	172	1	22	23	74
40	HNEDJ57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	50	1813	1	1184	2	2	173	1	1	2	333

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
41	HNTME13	97901 02/26/97 209047 05/15/97	pSport1	51	2070	74	2070	142	142	174	1	20	21	195
41	HNTME13	97901 02/26/97 209047 05/15/97	pSport1	109	2003	15	1957	68	68	232	1	22	23	300
42	HSXBI25	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	52	1426	1	1426	158	158	175	1	25	26	264
42	HSXBI25	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	110	1320	80	1311	41	41	233	1	29	30	312
43	HSXCK41	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	53	1720	1	1720	161	161	176	1	22	23	137
43	HSXCK41	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	111	1962	299	1962		566	234	1	33	34	47

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of AA
44	HE8CJ26	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	54	1117	1	1107	218	218	177	1	25	26	177
44	HE8CJ26	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	112	1785	30	1087		225	235	1	23	24	33
45	HTTDS54	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	55	1903	1	1903	119	119	178	1	31	32	154
45	HTTDS54	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	113	1842	1	1832	80	80	236	1	36	37	312
46	HLHDY31	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	56	1869	133	1838	124	124	179	1	24	25	294
46	HLHDY31	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	114	1960	90	1960	165	165	237	1	24	25	295

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
47	HMCBP63	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	57	1259	320	1010	352	352	180	1	26	27	255
48	HEMGE83	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	58	1186	33	557	12	12	181	1	18	19	323
49	HHSDC22	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	59	428	1	304	172	172	182	1	34	35	46
50	HHSDZ57	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	60	501	1	501	40	40	183	1	62	63	92
50	HHSDZ57	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	115	536	73	536	73	73	238	1	22	23	91
52	HMMAB12	97903 02/26/97 209049 05/15/97	pSport1	62	595	1	595	308	308	185	1	29	30	42

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
52	HMMAB12	97903 02/26/97 209049 05/15/97	pSport1	118	453	1	453	198	198	241	1	26	27	27
53	HSKDW02	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	63	1478	40	1436	176	176	186	1	39	40	58
53	HSKDW02	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	119	2016	211	1957	317	317	242	1	25	26	57
54	HETGL41	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	64	2033	1	2033	30	30	187	1	30	31	187
54	HETGL41	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	120	2136	110	2134	296	296	243	1	23	24	122
55	HODAZ50	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	65	440	1	440	1	1	188	1	26	27	145

Gene No.	cDNA Clone ID	ATCC Deposit No.: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
55	HODAZ50	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	121	219	1	219		1	244	1	10	11	72
56	HSDGE59	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	66	3301	349	1478	341	341	189	1	30	31	83
57	HE6ES13	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	67	1535	1	1535	331	331	190	1	26	27	57
57	HE6ES13	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	122	1686	239	1678		367	245	1	27	28	48
58	HSSEP68	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	68	1244	402	1244	57	57	191	1	30	31	310
58	HSSEP68	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	123	1211	1	1211	80	80	246	1	30	31	338

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
58	HSSEP68	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	124	1804	402	1526	501	501	247	1			17
59	HRDEV41	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	69	1292	1	1278	70	70	192	1	28	29	317
59	HRDEV41	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	125	1282	31	1088	70	70	248	1	21	22	338
60	HILCJ01	97903 02/26/97 209049 05/15/97	pBluescript SK-	70	1031	498	1031	536	536	193	1	30	31	52
61	HSATP28	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	71	855	178	855	187	187	194	1	28	29	41
62	HHFGL41	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	72	1274	58	1274	118	118	195	1	42	43	101

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
62	HHFGL41	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	126	1296	88	1237	133	133	249	1	39	40	95
63	HBJEM49	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	73	688	1	688	173	173	196	1	18	19	44
63	HBJEM49	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	127	737	1	737	174	174	250	1	20	21	78
64	HSLDJ95	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	74	1890	1	1890	112	112	197	1	21	22	354
64	HSLDJ95	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	128	1925	1	1829	87	87	251	1	23	24	353
65	HSREG44	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	75	1133	408	1133	531	531	198	1	18	19	73

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
66	HTXCT40	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	76	585	1	585	1	1	199	1	69	70	112
66	HTXCT40	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	129	2713	2023	2713	2133	2133	252	1	39	40	108
67	HRGDF73	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	77	577	1	577	51	51	200	1	23	24	122
68	HRDBF52	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	78	2278	1458	1935	25	25	201	1	23	24	314
68	HRDBF52	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	130	1011	479	1011	701	701	253	1	20	21	44
68	HKMND45	97976 04/04/97	pBluescript	131	2278	1	1929	25	25	254	1	27	28	314
69	HPEBD70	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	79	1143	601	1097	95	95	202	1	6	7	235

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of AA of Signal Pep	AA SEQ ID NO: Y	First AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of OR F
69	HPEBD70	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	132	1088	535	1043	588	588	255	1	27	28	52
70	HMCAB89	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	80	557	1	557	132	132	203	1	25	26	92

Table 1 summarizes the information corresponding to each "Gene No." described above. The nucleotide sequence identified as "NT SEQ ID NO:X" was assembled from partially homologous ("overlapping") sequences obtained from the "cDNA clone ID" identified in Table 1 and, in some cases, from additional related DNA clones. The overlapping sequences were assembled into a single contiguous sequence of high redundancy (usually three to five overlapping sequences at each nucleotide position), resulting in a final sequence identified as SEQ ID NO:X.

The cDNA Clone ID was deposited on the date and given the corresponding deposit number listed in "ATCC Deposit No:Z and Date." Some of the deposits contain multiple different clones corresponding to the same gene. "Vector" refers to the type of vector contained in the cDNA Clone ID.

"Total NT Seq." refers to the total number of nucleotides in the contig identified by "Gene No." The deposited clone may contain all or most of these sequences, reflected by the nucleotide position indicated as "5' NT of Clone Seq." and the "3' NT of Clone Seq." of SEQ ID NO:X. The nucleotide position of SEQ ID NO:X of the putative start codon (methionine) is identified as "5' NT of Start Codon." Similarly, the nucleotide position of SEQ ID NO:X of the predicted signal sequence is identified as "5' NT of First AA of Signal Pep."

The translated amino acid sequence, beginning with the methionine, is identified as "AA SEQ ID NO:Y," although other reading frames can also be easily translated using known molecular biology techniques. The polypeptides produced by these alternative open reading frames are specifically contemplated by the present invention.

The first and last amino acid position of SEQ ID NO:Y of the predicted signal peptide is identified as "First AA of Sig Pep" and "Last AA of Sig Pep." The predicted first amino acid position of SEQ ID NO:Y of the secreted portion is identified as "Predicted First AA of Secreted Portion." Finally, the amino acid position of SEQ ID NO:Y of the last amino acid in the open reading frame is identified as "Last AA of ORF."

SEQ ID NO:X and the translated SEQ ID NO:Y are sufficiently accurate and otherwise suitable for a variety of uses well known in the art and described further below. For instance, SEQ ID NO:X is useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO:X or the cDNA contained in the deposited clone. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO:Y may be used to generate antibodies which bind specifically to the secreted proteins encoded by the cDNA clones identified in Table 1.

Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

Accordingly, for those applications requiring precision in the nucleotide sequence or the amino acid sequence, the present invention provides not only the generated nucleotide sequence identified as SEQ ID NO:X and the predicted translated amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA containing a human cDNA of the invention deposited with the ATCC, as set forth in Table 1. The nucleotide sequence of each deposited clone can readily be determined by sequencing the deposited clone in accordance with known methods. The predicted amino acid sequence can then be verified from such deposits. Moreover, the amino acid sequence of the protein encoded by a particular clone can also be directly determined by peptide sequencing or by expressing the protein in a suitable host cell containing the deposited human cDNA, collecting the protein, and determining its sequence.

The present invention also relates to the genes corresponding to SEQ ID NO:X, SEQ ID NO:Y, or the deposited clone. The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

Also provided in the present invention are species homologs. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source for the desired homologue.

The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood in the art.

The polypeptides may be in the form of the secreted protein, including the mature form, or may be a part of a larger protein, such as a fusion protein (see below).

It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple histidine residues, or an additional sequence for stability during recombinant production.

- 5 The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of a polypeptide, including the secreted polypeptide, can be substantially purified by the one-step method described in Smith and Johnson, *Gene* 67:31-40 (1988). Polypeptides of the invention also can be purified from natural or recombinant sources
- 10 using antibodies of the invention raised against the secreted protein in methods which are well known in the art.

Signal Sequences

- 15 Methods for predicting whether a protein has a signal sequence, as well as the cleavage point for that sequence, are available. For instance, the method of McGeoch, *Virus Res.* 3:271-286 (1985), uses the information from a short N-terminal charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje, *Nucleic Acids Res.* 14:4683-4690 (1986) uses the information from the residues surrounding the cleavage site, typically residues -13 to +2, where +1
- 20 indicates the amino terminus of the secreted protein. The accuracy of predicting the cleavage points of known mammalian secretory proteins for each of these methods is in the range of 75-80%. (von Heinje, *supra*.) However, the two methods do not always produce the same predicted cleavage point(s) for a given protein.

- 25 In the present case, the deduced amino acid sequence of the secreted polypeptide was analyzed by a computer program called SignalP (Henrik Nielsen et al., *Protein Engineering* 10:1-6 (1997)), which predicts the cellular location of a protein based on the amino acid sequence. As part of this computational prediction of localization, the methods of McGeoch and von Heinje are incorporated. The analysis of the amino acid sequences of the secreted proteins described herein by this program provided the results
- 30 shown in Table 1.

- As one of ordinary skill would appreciate, however, cleavage sites sometimes vary from organism to organism and cannot be predicted with absolute certainty. Accordingly, the present invention provides secreted polypeptides having a sequence shown in SEQ ID NO:Y which have an N-terminus beginning within 5 residues (i.e., +
- 35 or - 5 residues) of the predicted cleavage point. Similarly, it is also recognized that in some cases, cleavage of the signal sequence from a secreted protein is not entirely

uniform, resulting in more than one secreted species. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

Moreover, the signal sequence identified by the above analysis may not necessarily predict the naturally occurring signal sequence. For example, the naturally occurring signal sequence may be further upstream from the predicted signal sequence. However, it is likely that the predicted signal sequence will be capable of directing the secreted protein to the ER. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

10 Polynucleotide and Polypeptide Variants

"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

"Identity" per se has an art-recognized meaning and can be calculated using published techniques. (See, e.g.: (COMPUTATIONAL MOLECULAR BIOLOGY, Lesk, A.M., ed., Oxford University Press, New York, (1988); BIOCOMPUTING: INFORMATICS AND GENOME PROJECTS, Smith, D.W., ed., Academic Press, New York, (1993); COMPUTER ANALYSIS OF SEQUENCE DATA, PART I, Griffin, A.M., and Griffin, H.G., eds., Humana Press, New Jersey, (1994); SEQUENCE ANALYSIS IN MOLECULAR BIOLOGY, von Heinje, G., Academic Press, (1987); and SEQUENCE ANALYSIS PRIMER, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, (1991).) While there exists a number of methods to measure identity between two polynucleotide or polypeptide sequences, the term "identity" is well known to skilled artisans. (Carillo, H., and Lipton, D., SIAM J Applied Math 48:1073 (1988).) Methods commonly employed to determine identity or similarity between two sequences include, but are not limited to, those disclosed in "Guide to Huge Computers," Martin J. Bishop, ed., Academic Press, San Diego, (1994), and Carillo, H., and Lipton, D., SIAM J Applied Math 48:1073 (1988). Methods for aligning polynucleotides or polypeptides are codified in computer programs, including the GCG program package (Devereux, J., et al., Nucleic Acids Research (1984) 12(1):387 (1984)), BLASTP, BLASTN, FASTA (Atschul, S.F. et al., J. Molec. Biol. 215:403 (1990)), Bestfit program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, 575 Science Drive, Madison, WI 53711 (using the local homology algorithm of Smith and Waterman, Advances in Applied Mathematics 2:482-489 (1981)).

When using any of the sequence alignment programs to determine whether a particular sequence is, for instance, 95% identical to a reference sequence, the parameters are set so that the percentage of identity is calculated over the full length of the reference polynucleotide and that gaps in identity of up to 5% of the total number of nucleotides in the reference polynucleotide are allowed.

A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. 6:237-245 (1990).) The term "sequence" includes nucleotide and amino acid sequences. In a sequence alignment the query and subject sequences are either both nucleotide sequences or both amino acid sequences. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB search of a DNA sequence to calculate percent identity are: Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization Group Length=0, and Cutoff Score=1, Gap Penalty=5, Gap Size Penalty 0.05, and Window Size=500 or query sequence length in nucleotide bases, whichever is shorter. Preferred parameters employed to calculate percent identity and similarity of an amino acid alignment are: Matrix=PAM 150, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty=0.05, and Window Size=500 or query sequence length in amino acid residues, whichever is shorter.

As an illustration, a polynucleotide having a nucleotide sequence of at least 95% "identity" to a sequence contained in SEQ ID NO:X or the cDNA contained in the deposited clone, means that the polynucleotide is identical to a sequence contained in SEQ ID NO:X or the cDNA except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the total length (not just within a given 100 nucleotide stretch). In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to SEQ ID NO:X or the deposited clone, up to 5% of the nucleotides in the sequence contained in SEQ ID NO:X or the cDNA can be deleted, inserted, or substituted with other nucleotides. These changes may occur anywhere throughout the polynucleotide.

Further embodiments of the present invention include polynucleotides having at least 85% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to a sequence contained in SEQ ID NO:X or the cDNA contained in the deposited clone. Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the polynucleotides having at least 85%, 90%, 95%, 96%, 97%, 98%, or 99% identity

will encode a polypeptide identical to an amino acid sequence contained in SEQ ID NO:Y or the expressed protein produced by the deposited clone.

Similarly, by a polypeptide having an amino acid sequence having at least, for example, 95% "identity" to a reference polypeptide, is intended that the amino acid
5 sequence of the polypeptide is identical to the reference polypeptide except that the polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the total length of the reference polypeptide. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a reference amino acid sequence, up to 5% of the amino acid residues in the reference sequence may be
10 deleted or substituted with another amino acid, or a number of amino acids up to 5% of the total amino acid residues in the reference sequence may be inserted into the reference sequence. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the
15 reference sequence or in one or more contiguous groups within the reference sequence.

Further embodiments of the present invention include polypeptides having at least 80% identity, more preferably at least 85% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to an amino acid sequence contained in SEQ ID NO:Y or the expressed protein produced by
20 the deposited clone. Preferably, the above polypeptides should exhibit at least one biological activity of the protein.

In a preferred embodiment, polypeptides of the present invention include polypeptides having at least 90% similarity, more preferably at least 95% similarity, and still more preferably at least 96%, 97%, 98%, or 99% similarity to an amino acid
25 sequence contained in SEQ ID NO:Y or the expressed protein produced by the deposited clone.

The variants may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or
30 activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in
35 the human mRNA to those preferred by a bacterial host such as *E. coli*).

Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an

organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level.

Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

5 Using known methods of protein engineering and recombinant DNA technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein without substantial loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988
10 (1993), reported variant KGF proteins having heparin binding activity even after deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).)

 Moreover, ample evidence demonstrates that variants often retain a biological
15 activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1a. They used random mutagenesis to generate over 3,500 individual IL-1a mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible
20 amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." (See, Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

25 Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form
30 are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

 Thus, the invention further includes polypeptide variants which show
35 substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make

phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

5 The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid
10 substitution could be modified while still maintaining biological activity of the protein.

The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham
15 and Wells, Science 244:1081-1085 (1989).) The resulting mutant molecules can then be tested for biological activity.

As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the
20 protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues
25 Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues,
30 where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino
35 acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

Polynucleotide and Polypeptide Fragments

10 In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in the deposited clone or shown in SEQ ID NO:X. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in
15 length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in the deposited clone or the nucleotide sequence shown in SEQ ID NO:X. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, 500, 600, 2000 nucleotides) are preferred.

20 Moreover, representative examples of polynucleotide fragments of the invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, and 701 to the end of SEQ ID NO:X or the cDNA contained in the deposited clone. In this context "about" includes the particularly
25 recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has biological activity.

In the present invention, a "polypeptide fragment" refers to a short amino acid sequence contained in SEQ ID NO:Y or encoded by the cDNA contained in the
30 deposited clone. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, and 161 to the end of the coding
35 region. Moreover, polypeptide fragments can be about 20, 30, 40, 50, 60, 70, 80, 90, 100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about"

includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or
5 the mature form having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any
10 combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alpha-
15 helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions. Polypeptide fragments of SEQ ID NO:Y falling within conserved domains are
20 specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the
25 fragments may include an improved desired activity, or a decreased undesirable activity.

Epitopes & Antibodies

In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred
30 embodiment of the present invention relates to a polypeptide fragment comprising an epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA
35 81:3998- 4002 (1983).)

Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

5 In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

10 Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if
15 it is long enough (at least about 25 amino acids), without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

20 As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')₂ fragments) which are capable of specifically binding to protein. Fab and F(ab')₂ fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding than an intact antibody. (Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred,
25 as well as the products of a FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

Fusion Proteins

30 Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular
35 locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

5 Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the
10 polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

 Moreover, polypeptides of the present invention, including fragments, and specifically epitopes, can be combined with parts of the constant domain of
15 immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life in vivo. One reported example describes chimeric proteins consisting of the first two domains of the human CD4- polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86
20 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) can also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., J. Biochem. 270:3958-3964 (1995).)

 Similarly, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion
25 proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. (EP-A 0232 262.) Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified,
30 would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D. Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol.
35 Chem. 270:9459-9471 (1995).)

 Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a peptide which facilitates purification of the fused polypeptide. In

preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).)

Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the claimed invention.

Vectors, Host Cells, and Protein Production

The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged in vitro using an appropriate packaging cell line and then transduced into host cells.

The polynucleotide insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination, and, in the transcribed region, a ribosome binding site for translation. The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin resistance for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance genes for culturing in E. coli and other bacteria. Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as E. coli, Streptomyces and Salmonella typhimurium cells; fungal cells, such as yeast cells; insect cells such as Drosophila S2 and Spodoptera Sf9 cells; animal cells such as CHO, COS,

293, and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan.

Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis et al., Basic Methods In Molecular Biology (1986). It is specifically contemplated that the polypeptides of the present invention may in fact be expressed by a host cell lacking a recombinant vector.

A polypeptide of this invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification.

Polypeptides of the present invention, and preferably the secreted form, can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any protein after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient, depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

Uses of the Polynucleotides

Each of the polynucleotides identified herein can be used in numerous ways as reagents. The following description should be considered exemplary and utilizes
5 known techniques.

The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat polymorphisms), are presently available. Each polynucleotide of the present invention
10 can be used as a chromosome marker.

Briefly, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can be selected using computer analysis so that primers do not span more than one predicted exon in the genomic DNA. These primers are then used for PCR screening of somatic
15 cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the SEQ ID NO:X will yield an amplified fragment.

Similarly, somatic hybrids provide a rapid method of PCR mapping the polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can
20 be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flow-sorted chromosomes, and preselection by hybridization to construct chromosome specific-cDNA libraries.

Precise chromosomal location of the polynucleotides can also be achieved using
25 fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides 2,000-4,000 bp are preferred. For a review of this technique, see Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York (1988).

For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for marking multiple sites and/or multiple chromosomes). Preferred polynucleotides correspond to the noncoding regions of the cDNAs because the coding sequences are more likely conserved within gene families, thus increasing the chance of cross
30 hybridization during chromosomal mapping.

Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage

analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library) .) Assuming 1 megabase mapping resolution and one gene per
5 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential causative genes.

Thus, once coinheritance is established, differences in the polynucleotide and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or
10 translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the
15 mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using polynucleotides of the present invention. Any of these alterations (altered expression,
20 chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker.

In addition to the foregoing, a polynucleotide can be used to control gene expression through triple helix formation or antisense DNA or RNA. Both methods rely on binding of the polynucleotide to DNA or RNA. For these techniques, preferred
25 polynucleotides are usually 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression, CRC
30 Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques are effective in model systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease.

35 Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the

present invention offers means of targeting such genetic defects in a highly accurate manner. Another gene can insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell.

5 The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for RFLP.

10 The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

20 Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992).) Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

30 There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers specific to particular tissue prepared from the sequences of the present invention. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for contamination.

In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

Uses of the Polypeptides

Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

A polypeptide of the present invention can be used to assay protein levels in a biological sample using antibody-based techniques. For example, protein expression in tissues can be studied with classical immunohistological methods. (Jalkanen, M., et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell . Biol. 105:3087-3096 (1987).) Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (125I, 121I), carbon (14C), sulfur (35S), tritium (3H), indium (112In), and technetium (99mTc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

In addition to assaying secreted protein levels in a biological sample, proteins can also be detected in vivo by imaging. Antibody labels or markers for in vivo imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 131I, 112In, 99mTc), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously, or intraperitoneally) into the mammal. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20

millicuries of ^{99m}Tc . The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. In vivo tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Tumor Imaging: The Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982).)

Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression of a polypeptide of the present invention in cells or body fluid of an individual; (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a disorder.

Moreover, polypeptides of the present invention can be used to treat disease. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B), to inhibit the activity of a polypeptide (e.g., an oncogene), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth).

Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease. For example, administration of an antibody directed to a polypeptide of the present invention can bind and reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

At the very least, the polypeptides of the present invention could be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the following biological activities.

Biological Activities

The polynucleotides and polypeptides of the present invention can be used in assays to test for one or more biological activities. If these polynucleotides and polypeptides do exhibit activity in a particular assay, it is likely that these molecules

may be involved in the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides could be used to treat the associated disease.

Immune Activity

5 A polypeptide or polynucleotide of the present invention may be useful in treating deficiencies or disorders of the immune system, by activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells
10 from pluripotent stem cells. The etiology of these immune deficiencies or disorders may be genetic, somatic, such as cancer or some autoimmune disorders, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, a polynucleotide or polypeptide of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

15 A polynucleotide or polypeptide of the present invention may be useful in treating or detecting deficiencies or disorders of hematopoietic cells. A polypeptide or polynucleotide of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat those disorders associated with a decrease in certain (or many) types hematopoietic
20 cells. Examples of immunologic deficiency syndromes include, but are not limited to: blood protein disorders (e.g. agammaglobulinemia, dysgammaglobulinemia), ataxia telangiectasia, common variable immunodeficiency, Digeorge Syndrome, HIV infection, HTLV-BLV infection, leukocyte adhesion deficiency syndrome, lymphopenia, phagocyte bactericidal dysfunction, severe combined immunodeficiency
25 (SCIDs), Wiskott-Aldrich Disorder, anemia, thrombocytopenia, or hemoglobinuria.

 Moreover, a polypeptide or polynucleotide of the present invention could also be used to modulate hemostatic (the stopping of bleeding) or thrombolytic activity (clot formation). For example, by increasing hemostatic or thrombolytic activity, a polynucleotide or polypeptide of the present invention could be used to treat blood
30 coagulation disorders (e.g., afibrinogenemia, factor deficiencies), blood platelet disorders (e.g. thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, a polynucleotide or polypeptide of the present invention that can decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment of heart attacks
35 (infarction), strokes, or scarring.

 A polynucleotide or polypeptide of the present invention may also be useful in treating or detecting autoimmune disorders. Many autoimmune disorders result from

inappropriate recognition of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing autoimmune disorders.

Examples of autoimmune disorders that can be treated or detected by the present invention include, but are not limited to: Addison's Disease, hemolytic anemia, antiphospholipid syndrome, rheumatoid arthritis, dermatitis, allergic encephalomyelitis, glomerulonephritis, Goodpasture's Syndrome, Graves' Disease, Multiple Sclerosis, Myasthenia Gravis, Neuritis, Ophthalmia, Bullous Pemphigoid, Pemphigus, Polyendocrinopathies, Purpura, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Thyroiditis, Systemic Lupus Erythematosus, Autoimmune Pulmonary Inflammation, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disease.

Similarly, allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated by a polypeptide or polynucleotide of the present invention. Moreover, these molecules can be used to treat anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

A polynucleotide or polypeptide of the present invention may also be used to treat and/or prevent organ rejection or graft-versus-host disease (GVHD). Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. The administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing organ rejection or GVHD.

Similarly, a polypeptide or polynucleotide of the present invention may also be used to modulate inflammation. For example, the polypeptide or polynucleotide may inhibit the proliferation and differentiation of cells involved in an inflammatory response. These molecules can be used to treat inflammatory conditions, both chronic and acute conditions, including inflammation associated with infection (e.g., septic shock, sepsis, or systemic inflammatory response syndrome (SIRS)), ischemia-reperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel disease, Crohn's disease, or resulting from over production of cytokines (e.g., TNF or IL-1.)

Hyperproliferative Disorders

A polypeptide or polynucleotide can be used to treat or detect hyperproliferative disorders, including neoplasms. A polypeptide or polynucleotide of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, a polypeptide or polynucleotide of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative disorders can be treated. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating hyperproliferative disorders, such as a chemotherapeutic agent.

Examples of hyperproliferative disorders that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but are not limited to neoplasms located in the: abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvic, skin, soft tissue, spleen, thoracic, and urogenital.

Similarly, other hyperproliferative disorders can also be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome, Waldenstrom's Macroglobulinemia, Gaucher's Disease, histiocytosis, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

Infectious Disease

A polypeptide or polynucleotide of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, the polypeptide or polynucleotide of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

- Viruses are one example of an infectious agent that can cause disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of viruses, include, but are not limited to the following DNA and RNA viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus,
- 5 Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza), Papovaviridae, Parvoviridae, Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g.,
- 10 Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis, bronchiolitis, encephalitis, eye infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), meningitis, opportunistic infections (e.g., AIDS),
- 15 pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.
- 20 Similarly, bacterial or fungal agents that can cause disease or symptoms and that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following Gram-Negative and Gram-positive bacterial families and fungi: Actinomycetales (e.g., Corynebacterium, Mycobacterium, Norcardia), Aspergillosis, Bacillaceae (e.g., Anthrax, Clostridium), Bacteroidaceae,
- 25 Blastomycosis, Bordetella, Borrelia, Brucellosis, Candidiasis, Campylobacter, Coccidioidomycosis, Cryptococcosis, Dermatocycoses, Enterobacteriaceae (Klebsiella, Salmonella, Serratia, Yersinia), Erysipelothrix, Helicobacter, Legionellosis, Leptospirosis, Listeria, Mycoplasmatales, Neisseriaceae (e.g., Acinetobacter, Gonorrhea, Meningococcal), Pasteurellaceae Infections (e.g., Actinobacillus,
- 30 Heamophilus, Pasteurella), Pseudomonas, Rickettsiaceae, Chlamydiaceae, Syphilis, and Staphylococcal. These bacterial or fungal families can cause the following diseases or symptoms, including, but not limited to: bacteremia, endocarditis, eye infections (conjunctivitis, tuberculosis, uveitis), gingivitis, opportunistic infections (e.g., AIDS related infections), paronychia, prosthesis-related infections, Reiter's Disease,
- 35 respiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, Dysentery, Paratyphoid Fever, food poisoning, Typhoid, pneumonia, Gonorrhea, meningitis, Chlamydia, Syphilis, Diphtheria.

Leprosy, Paratuberculosis, Tuberculosis, Lupus, Botulism, gangrene, tetanus, impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections. A polypeptide or polynucleotide of the present invention can be used to treat or detect
5 any of these symptoms or diseases.

Moreover, parasitic agents causing disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following families: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis, Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis,
10 Leishmaniasis, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas. These parasites can cause a variety of diseases or symptoms, including, but not limited to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), Malaria, pregnancy complications, and toxoplasmosis. A polypeptide or polynucleotide
15 of the present invention can be used to treat or detect any of these symptoms or diseases.

Preferably, treatment using a polypeptide or polynucleotide of the present invention could either be by administering an effective amount of a polypeptide to the patient, or by removing cells from the patient, supplying the cells with a polynucleotide
20 of the present invention, and returning the engineered cells to the patient (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease.

Regeneration

25 A polynucleotide or polypeptide of the present invention can be used to differentiate, proliferate, and attract cells, leading to the regeneration of tissues. (See, Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteoarthritis, periodontal
30 disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion injury, or systemic cytokine damage.

Tissues that could be regenerated using the present invention include organs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac), vascular (including vascular endothelium), nervous, hematopoietic, and
35 skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs without or decreased scarring. Regeneration also may include angiogenesis.

Moreover, a polynucleotide or polypeptide of the present invention may increase regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. A polynucleotide or polypeptide of the present invention could also be used prophylactically in an effort to avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel syndrome, and other tendon or ligament defects. A further example of tissue regeneration of non-healing wounds includes pressure ulcers, ulcers associated with vascular insufficiency, surgical, and traumatic wounds.

Similarly, nerve and brain tissue could also be regenerated by using a polynucleotide or polypeptide of the present invention to proliferate and differentiate nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and stroke). Specifically, diseases associated with peripheral nerve injuries, peripheral neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome), could all be treated using the polynucleotide or polypeptide of the present invention.

Chemotaxis

A polynucleotide or polypeptide of the present invention may have chemotaxis activity. A chemotactic molecule attracts or mobilizes cells (e.g., monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells) to a particular site in the body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

A polynucleotide or polypeptide of the present invention may increase chemotactic activity of particular cells. These chemotactic molecules can then be used to treat inflammation, infection, hyperproliferative disorders, or any immune system disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotactic molecules can be used to treat wounds and other trauma to tissues by attracting immune cells to the injured location. Chemotactic molecules of the present invention can also attract fibroblasts, which can be used to treat wounds.

It is also contemplated that a polynucleotide or polypeptide of the present invention may inhibit chemotactic activity. These molecules could also be used to treat

disorders. Thus, a polynucleotide or polypeptide of the present invention could be used as an inhibitor of chemotaxis.

Binding Activity

5 A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or
10 small molecules.

Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural
15 receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide, either as a secreted protein or on the cell
20 membrane. Preferred cells include cells from mammals, yeast, *Drosophila*, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the molecule to observe binding, stimulation, or inhibition of activity of either the polypeptide or the molecule.

25 The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

Alternatively, the assay can be carried out using cell-free preparations.
30 polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a standard.

35 Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The

antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptide from suitably manipulated cells or tissues.

Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

Other Activities

A polypeptide or polynucleotide of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

A polypeptide or polynucleotide of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide or polynucleotide of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

A polypeptide or polynucleotide of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, circadian rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity), hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

A polypeptide or polynucleotide of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional components.

Other Preferred Embodiments

Other preferred embodiments of the claimed invention include an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1.

Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Clone Sequence and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Start Codon and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Similarly preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 150 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

Further preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 500 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of SEQ ID NO:X beginning with the nucleotide at about the position of the 5' Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NO:X.

Also preferred is an isolated nucleic acid molecule which hybridizes under stringent hybridization conditions to a nucleic acid molecule, wherein said nucleic acid molecule which hybridizes does not hybridize under stringent hybridization conditions to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or of only T residues.

Also preferred is a composition of matter comprising a DNA molecule which comprises a human cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the material deposited with the American Type Culture Collection and given the ATCC Deposit Number shown in Table 1 for said cDNA Clone Identifier.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in the nucleotide sequence of a human cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the deposit given the ATCC Deposit Number shown in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of the complete open reading frame sequence encoded by said human cDNA clone.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is a method for detecting in a biological sample a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method comprises a step of comparing a nucleotide sequence of at least one nucleic acid molecule in said sample with a sequence selected from said group and determining

whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

10 A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any
15 integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide
20 sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample
25 obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide
30 sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least
35 two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1.

Also preferred is a polypeptide, wherein said sequence of contiguous amino acids is included in the amino acid sequence of SEQ ID NO:Y in the range of positions beginning with the residue at about the position of the First Amino Acid of the Secreted Portion and ending with the residue at about the Last Amino Acid of the Open Reading Frame as set forth for SEQ ID NO:Y in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a secreted portion of the secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the

amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at
5 least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at
10 least 95% identical to the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is an isolated antibody which binds specifically to a
15 polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in
20 the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1;
25 and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method comprises a step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group and determining
30 whether the sequence of said polypeptide molecule in said sample is at least 90% identical to said sequence of at least 10 contiguous amino acids.

Also preferred is the above method wherein said step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of
35 polypeptides in said sample to an antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an

amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

5 Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

Also preferred is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting polypeptide molecules in
10 said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained
15 in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is the above method for identifying the species, tissue or cell type of a biological sample, which method comprises a step of detecting polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a
20 sequence of at least 10 contiguous amino acids in a sequence selected from the above group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample
25 obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid
30 sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

35 Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding a polypeptide wherein said polypeptide comprises an amino acid sequence that is at least

90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said nucleotide sequence encoding a polypeptide has been optimized for expression of said polypeptide in a prokaryotic host.

Also preferred is an isolated nucleic acid molecule, wherein said polypeptide comprises an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is a method of making a recombinant vector comprising inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is the recombinant vector produced by this method. Also preferred is a method of making a recombinant host cell comprising introducing the vector into a host cell, as well as the recombinant host cell produced by this method.

Also preferred is a method of making an isolated polypeptide comprising culturing this recombinant host cell under conditions such that said polypeptide is expressed and recovering said polypeptide. Also preferred is this method of making an isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said polypeptide is a secreted portion of a human secreted protein comprising an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y beginning with the residue at the position of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y wherein Y is an integer set forth in Table 1 and said position of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y is defined in Table 1; and an amino acid sequence of a secreted portion of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The isolated polypeptide produced by this method is also preferred.

Also preferred is a method of treatment of an individual in need of an increased level of a secreted protein activity, which method comprises administering to such an individual a pharmaceutical composition comprising an amount of an isolated

polypeptide, polynucleotide, or antibody of the claimed invention effective to increase the level of said protein activity in said individual.

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

Examples

Example 1: Isolation of a Selected cDNA Clone From the Deposited Sample

Each cDNA clone in a cited ATCC deposit is contained in a plasmid vector. Table 1 identifies the vectors used to construct the cDNA library from which each clone was isolated. In many cases, the vector used to construct the library is a phage vector from which a plasmid has been excised. The table immediately below correlates the related plasmid for each phage vector used in constructing the cDNA library. For example, where a particular clone is identified in Table 1 as being isolated in the vector "Lambda Zap," the corresponding deposited clone is in "pBluescript."

	<u>Vector Used to Construct Library</u>	<u>Corresponding Deposited Plasmid</u>
	Lambda Zap	pBluescript (pBS)
20	Uni-Zap XR	pBluescript (pBS)
	Zap Express	pBK
	lafmid BA	plafmid BA
	pSport I	pSport I
	pCMVSPORT 2.0	pCMVSPORT 2.0
25	pCMVSPORT 3.0	pCMVSPORT 3.0
	pCR [®] 2.1	pCR [®] 2.1

Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res. 16:7583-7600 (1988); Altting-Mees, M. A. and Short, J. M., Nucleic Acids Res. 17:9494 (1989)) and pBK (Altting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1 Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS. The S and K refers to the orientation of the polylinker to the T7 and T3 primer sequences which flank the polylinker region ("S" is for SacI and "K" is for KpnI which

are the first sites on each respective end of the linker). "+" or "-" refer to the orientation of the f1 origin of replication ("ori"), such that in one orientation, single stranded rescue initiated from the f1 ori generates sense strand DNA and in the other, antisense.

Vectors pSport1, pCMVSPORT 2.0 and pCMVSPORT 3.0, were obtained from
5 Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gruber, C. E., et al., Focus 15:59 (1993).) Vector lafmid BA (Bento Soares, Columbia University, NY) contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1
10 Blue. Vector pCR[®]2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark, J. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9: (1991).) Preferably, a polynucleotide of the present invention does not comprise the
15 phage vector sequences identified for the particular clone in Table 1, as well as the corresponding plasmid vector sequences designated above.

The deposited material in the sample assigned the ATCC Deposit Number cited in Table 1 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing
20 the same ATCC Deposit Number contain at least a plasmid for each cDNA clone identified in Table 1. Typically, each ATCC deposit sample cited in Table 1 comprises a mixture of approximately equal amounts (by weight) of about 50 plasmid DNAs, each containing a different cDNA clone; but such a deposit sample may include plasmids for more or less than 50 cDNA clones, up to about 500 cDNA clones.

25 Two approaches can be used to isolate a particular clone from the deposited sample of plasmid DNAs cited for that clone in Table 1. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to SEQ ID NO:X.

Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized
30 using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with ³²P-γ-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).) The plasmid mixture is transformed into a suitable host, as indicated above (such as
35 XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above.

The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., *Molecular Cloning: A Laboratory*
5 *Manual*, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1.93 to 1.104), or other techniques known to those of skill in the art.

Alternatively, two primers of 17-20 nucleotides derived from both ends of the SEQ ID NO:X (i.e., within the region of SEQ ID NO:X bounded by the 5' NT and the 3' NT of the clone defined in Table 1) are synthesized and used to amplify the desired
10 cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 µl of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation
15 at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the art. For instance, a method similar to 5' RACE is available for generating the
25 missing 5' end of a desired full-length transcript. (Fromont-Racine et al., *Nucleic Acids Res.* 21(7):1683-1684 (1993).)

Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to
30 a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.

This above method starts with total RNA isolated from the desired source, although poly-A⁺ RNA can be used. The RNA preparation can then be treated with
35 phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA which may interfere with the later RNA ligase step. The phosphatase should then

be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.

5 This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that
10 the 5' end sequence belongs to the desired gene.

Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

15 A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the cDNA sequence corresponding to SEQ ID NO:X., according to the method described in Example 1. (See also, Sambrook.)

Example 3: Tissue Distribution of Polypeptide

20 Tissue distribution of mRNA expression of polynucleotides of the present invention is determined using protocols for Northern blot analysis, described by, among others, Sambrook et al. For example, a cDNA probe produced by the method described in Example 1 is labeled with P³² using the rediprime™ DNA labeling system (Amersham Life Science), according to manufacturer's instructions. After labeling, the probe is purified using CHROMA SPIN-100™ column (Clontech Laboratories, Inc.),
25 according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various human tissues for mRNA expression.

30 Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) (Clontech) are examined with the labeled probe using ExpressHyb™ hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are mounted and exposed to film at -70°C overnight, and the films developed according to standard procedures.

Example 4: Chromosomal Mapping of the Polynucleotides

35 An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This

primer set is then used in a polymerase chain reaction under the following set of conditions : 30 seconds, 95°C; 1 minute, 56°C; 1 minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual
5 chromosomes or chromosome fragments (Bios, Inc). The reactions is analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

10 **Example 5: Bacterial Expression of a Polypeptide**

A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as
15 BamHI and XbaI, at the 5' end of the primers in order to clone the amplified product into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (Amp^r), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site
20 (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses
25 the lacI repressor and also confers kanamycin resistance (Kan^r). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis.

Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml).
30 The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D.⁶⁰⁰) of between 0.4 and 0.6. IPTG (Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression.

Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is removed by centrifugation, and the supernatant containing the polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., *supra*).

Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins are eluted by the addition of 250 mM imidazole. Imidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

In addition to the above expression vector, the present invention further includes an expression vector comprising phage operator and promoter elements operatively linked to a polynucleotide of the present invention, called pHE4a. (ATCC Accession Number XXXXXX.) This vector contains: 1) a neomycinphosphotransferase gene as a selection marker, 2) an E. coli origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgarno sequence, and 6) the lactose operon repressor gene (*lacIq*). The origin of replication (*oriC*) is derived from pUC19 (LTI, Gaithersburg, MD). The promoter sequence and operator sequences are made synthetically.

DNA can be inserted into the pHEa by restricting the vector with NdeI and XbaI, BamHI, XhoI, or Asp718, running the restricted product on a gel, and isolating the larger fragment (the stuffer fragment should be about 310 base pairs). The DNA insert is generated according to the PCR protocol described in Example 1, using PCR primers having restriction sites for NdeI (5' primer) and XbaI, BamHI, XhoI, or

Asp718 (3' primer). The PCR insert is gel purified and restricted with compatible enzymes. The insert and vector are ligated according to standard protocols.

The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

5

Example 6: Purification of a Polypeptide from an Inclusion Body

The following alternative method can be used to purify a polypeptide expressed in *E. coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

10 Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50
15 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

 The cells are then lysed by passing the solution through a microfluidizer (Microfluidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by
20 centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

 The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the polypeptide containing supernatant is incubated at 4°C
25 overnight to allow further GuHCl extraction.

 Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing
30 for 12 hours prior to further purification steps.

 To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive

Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

5 Fractions containing the polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium
10 acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A_{280} monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

15 The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from Commaesie blue stained 16% SDS-PAGE gel when 5 μ g of purified protein is loaded. The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

20

Example 7: Cloning and Expression of a Polypeptide in a Baculovirus Expression System

In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide into a baculovirus to express a polypeptide. This expression vector contains the strong
25 polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak *Drosophila* promoter in the
30 same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

Many other baculovirus vectors can be used in place of the vector above, such
35 as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription,

translocation, secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow et al., Virology 170:31-39 (1989).

Specifically, the cDNA sequence contained in the deposited clone, including the
5 AUG initiation codon and the naturally associated leader sequence identified in Table 1, is amplified using the PCR protocol described in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the pA2 vector does not need a second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a baculovirus leader sequence, using the standard methods described in Summers et al.,
10 "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures," Texas Agricultural Experimental Station Bulletin No. 1555 (1987).

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

15 The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("GeneClean" BIO 101 Inc., La Jolla, Ca.).

The fragment and the dephosphorylated plasmid are ligated together with T4
20 DNA ligase. *E. coli* HB101 or other suitable *E. coli* hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA
25 sequencing.

Five µg of a plasmid containing the polynucleotide is co-transfected with 1.0 µg of a commercially available linearized baculovirus DNA ("BaculoGold™ baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One µg of
30 BaculoGold™ virus DNA and 5 µg of the plasmid are mixed in a sterile well of a microtiter plate containing 50 µl of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10 µl Lipofectin plus 90 µl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm
35 tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate

and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, *supra*. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 µl of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 µCi of ³⁵S-methionine and 5 µCi ³⁵S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

Example 8: Expression of a Polypeptide in Mammalian Cells

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLV I, HIV I and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSPORT 2.0, and pCMVSPORT 3.0. Mammalian host cells that could be used
5 include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable
10 marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transfected cells.

The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of
15 interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta. 1097:107-143 (1990); Page, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991).) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); Bebbington et al., Bio/Technology 10:169-175 (1992). Using these markers, the
20 mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the
25 expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No. 209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985).) Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the
30 cloning of the gene of interest. The vectors also contain the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by
35 procedures known in the art. The vector is then isolated from a 1% agarose gel.

A polynucleotide of the present invention is amplified according to the protocol outlined in Example 1. If the naturally occurring signal sequence is used to produce the

secreted protein, the vector does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

5 The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then
10 transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene is used for transfection. Five μ g of the expression plasmid pC6 is cotransfected with 0.5 μ g of the plasmid pSVneo using lipofectin (Felgner et al., *supra*). The plasmid pSV2-neo
15 contains a dominant selectable marker, the *neo* gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418.
20 After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μ M, 2 μ M, 5 μ M, 10 mM, 20 mM). The same
25 procedure is repeated until clones are obtained which grow at a concentration of 100 - 200 μ M. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

Example 9: Protein Fusions

30 The polypeptides of the present invention are preferably fused to other proteins. These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding protein facilitates purification. (See Example 5; see also EP A 394.827; Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and
35 albumin increases the halflife time in vivo. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the

activity of a fusion protein. Fusion proteins can also create chimeric molecules having more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which
5 outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an
10 expression vector, preferably a mammalian expression vector.

For example, if pC4 (Accession No. 209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated
15 by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

If the naturally occurring signal sequence is used to produce the secreted protein, pC4 does not need a second signal peptide. Alternatively, if the naturally
20 occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

Human IgG Fc region:

GGGATCCGGAGCCCAAATCTTCTGACAAAACTCACACATGCCCACCGTGCC
25 CAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCCTCTTCCCCCAAACC
CAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTCACATGCGTGGTGGT
GGACGTAAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACG
GCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAAC
AGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACTGGCTG
30 AATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAACCCCC
ATCGAGAAAACCATCTCCAAGCCAAAGGGCAGCCCCGAGAACCACAGGT
GTACACCCTGCCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTCAGCCT
GACCTGCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGTGGAGTGGGA
GAGCAATGGGCAGCCGGAGAACAACACTACAAGACCACGCCTCCCGTGCTGG
35 ACTCCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCA
GGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGC

ACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGAGTGC
GACGGCCGCGACTCTAGAGGAT (SEQ ID NO:1)

Example 10: Production of an Antibody from a Polypeptide

5 The antibodies of the present invention can be prepared by a variety of methods.
(See, Current Protocols, Chapter 2.) For example, cells expressing a polypeptide of
the present invention is administered to an animal to induce the production of sera
containing polyclonal antibodies. In a preferred method, a preparation of the secreted
protein is prepared and purified to render it substantially free of natural contaminants.
10 Such a preparation is then introduced into an animal in order to produce polyclonal
antisera of greater specific activity.

 In the most preferred method, the antibodies of the present invention are
monoclonal antibodies (or protein binding fragments thereof). Such monoclonal
antibodies can be prepared using hybridoma technology. (Köhler et al., Nature
15 256:495 (1975); Köhler et al., Eur. J. Immunol. 6:511 (1976); Köhler et al., Eur. J.
Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell
Hybridomas, Elsevier, N.Y., pp. 563-681 (1981).) In general, such procedures
involve immunizing an animal (preferably a mouse) with polypeptide or, more
preferably, with a secreted polypeptide-expressing cell. Such cells may be cultured in
20 any suitable tissue culture medium; however, it is preferable to culture cells in Earle's
modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at
about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about
1,000 U/ml of penicillin, and about 100 µg/ml of streptomycin.

 The splenocytes of such mice are extracted and fused with a suitable myeloma
25 cell line. Any suitable myeloma cell line may be employed in accordance with the
present invention; however, it is preferable to employ the parent myeloma cell line
(SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are
selectively maintained in HAT medium, and then cloned by limiting dilution as
described by Wands et al. (Gastroenterology 80:225-232 (1981).) The hybridoma cells
30 obtained through such a selection are then assayed to identify clones which secrete
antibodies capable of binding the polypeptide.

 Alternatively, additional antibodies capable of binding to the polypeptide can be
produced in a two-step procedure using anti-idiotypic antibodies. Such a method
makes use of the fact that antibodies are themselves antigens, and therefore, it is
35 possible to obtain an antibody which binds to a second antibody. In accordance with
this method, protein specific antibodies are used to immunize an animal, preferably a

mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the protein-specific antibody can be blocked by the polypeptide. Such antibodies comprise anti-idiotypic antibodies to the protein-specific antibody and
5 can be used to immunize an animal to induce formation of further protein-specific antibodies.

It will be appreciated that Fab and F(ab')₂ and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such fragments are typically produced by proteolytic cleavage, using enzymes such as papain
10 (to produce Fab fragments) or pepsin (to produce F(ab')₂ fragments). Alternatively, secreted protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry.

For in vivo use of antibodies in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using
15 genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art. (See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO
20 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985).)

Example 11: Production Of Secreted Protein For High-Throughput Screening Assays

25

The following protocol produces a supernatant containing a polypeptide to be tested. This supernatant can then be used in the Screening Assays described in Examples 13-20.

First, dilute Poly-D-Lysine (644 587 Boehringer-Mannheim) stock solution
30 (1mg/ml in PBS) 1:20 in PBS (w/o calcium or magnesium 17-516F Biowhittaker) for a working solution of 50ug/ml. Add 200 ul of this solution to each well (24 well plates) and incubate at RT for 20 minutes. Be sure to distribute the solution over each well (note: a 12-channel pipetter may be used with tips on every other channel). Aspirate off the Poly-D-Lysine solution and rinse with 1ml PBS (Phosphate Buffered Saline). The
35 PBS should remain in the well until just prior to plating the cells and plates may be poly-lysine coated in advance for up to two weeks.

Plate 293T cells (do not carry cells past P+20) at 2×10^5 cells/well in .5ml DMEM(Dulbecco's Modified Eagle Medium)(with 4.5 G/L glucose and L-glutamine (12-604F Biowhittaker))/10% heat inactivated FBS(14-503F Biowhittaker)/1x Penstrep(17-602E Biowhittaker). Let the cells grow overnight.

5 The next day, mix together in a sterile solution basin: 300 ul Lipofectamine (18324-012 Gibco/BRL) and 5ml Optimem I (31985070 Gibco/BRL)/96-well plate. With a small volume multi-channel pipetter, aliquot approximately 2ug of an expression vector containing a polynucleotide insert, produced by the methods described in Examples 8 or 9, into an appropriately labeled 96-well round bottom plate. With a
10 multi-channel pipetter, add 50ul of the Lipofectamine/Optimem I mixture to each well. Pipette up and down gently to mix. Incubate at RT 15-45 minutes. After about 20 minutes, use a multi-channel pipetter to add 150ul Optimem I to each well. As a control, one plate of vector DNA lacking an insert should be transfected with each set of transfections.

15 Preferably, the transfection should be performed by tag-teaming the following tasks. By tag-teaming, hands on time is cut in half, and the cells do not spend too much time on PBS. First, person A aspirates off the media from four 24-well plates of cells, and then person B rinses each well with .5-1ml PBS. Person A then aspirates off PBS rinse, and person B, using a 12-channel pipetter with tips on every other channel,
20 adds the 200ul of DNA/Lipofectamine/Optimem I complex to the odd wells first, then to the even wells, to each row on the 24-well plates. Incubate at 37°C for 6 hours.

 While cells are incubating, prepare appropriate media, either 1%BSA in DMEM with 1x penstrep, or CHO-5 media (see below) with 2mm glutamine and 1x penstrep. (BSA (81-068-3 Bayer) 100gm dissolved in 1L DMEM for a 10% BSA stock
25 solution). Filter the media and collect 50 ul for endotoxin assay in 15ml polystyrene conical.

 The transfection reaction is terminated, preferably by tag-teaming, at the end of the incubation period. Person A aspirates off the transfection media, while person B adds 1.5ml appropriate media to each well. Incubate at 37°C for 45 or 72 hours
30 depending on the media used: 1%BSA for 45 hours or CHO-5 for 72 hours.

 On day four, using a 300ul multichannel pipetter, aliquot 600ul in one 1ml deep well plate and the remaining supernatant into a 2ml deep well. The supernatants from each well can then be used in the assays described in Examples 13-20.

 It is specifically understood that when activity is obtained in any of the assays
35 described below using a supernatant, the activity originates from either the polypeptide directly (e.g., as a secreted protein) or by the polypeptide inducing expression of other

proteins, which are then secreted into the supernatant. Thus, the invention further provides a method of identifying the protein in the supernatant characterized by an activity in a particular assay.

5 *HGS-CHO-5 medium formulation:*

Inorganic Salts

CaCl ₂ (anhyd)	116.6 mg/L
CuSO ₄ ·5H ₂ O	0.00130
Fe(NO ₃) ₃ ·9H ₂ O	0.050
FeSO ₄ ·7H ₂ O	0.417
KCl	311.80
MgCl ₂	28.64
MgSO ₄	48.84
NaCl	6995.50
NaHCO ₃	2400.0
NaH ₂ PO ₄ ·H ₂ O	62.50
Na ₂ HPO ₄	71.02
ZnSO ₄ ·7H ₂ O	.4320

Lipids

Arachidonic Acid	.002 mg/L
Cholesterol	1.022
DL-alpha-Tocopherol-Acetate	.070
Linoleic Acid	0.0520
Linolenic Acid	0.010
Myristic Acid	0.010
Oleic Acid	0.010
Palmitric Acid	0.010
Palmitic Acid	0.010
Pluronic F-68	100
Stearic Acid	0.010
Tween 80	2.20

10 Carbon Source

D-Glucose	4551 mg/L
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Amino Acids

L- Alanine	130.85 mg/ml
L-Arginine-HCL	147.50
L-Asparagine-H ₂ O	7.50

L-Aspartic Acid	6.65
L-Cystine-2HCL-H ₂ O	29.56
L-Cystine-2HCL	31.29
L-Glutamic Acid	7.35
L-Glutamine	365.0
Glycine	18.75
L-Histidine-HCL-H ₂ O	52.48
L-Isoleucine	106.97
L-Leucine	111.45
L-Lysine HCL	163.75
L-Methionine	32.34
L-Phenylalanine	68.48
L-Proline	40.0
L-Serine	26.25
L-Threonine	101.05
L-Tryptophan	19.22
L-Tyrosine-2Na-2H ₂ O	91.79
L-Valine	99.65

Vitamins

Biotin	0.0035 mg/L
D-Ca Pantothenate	3.24
Choline Chloride	11.78
Folic Acid	4.65
i-Inositol	15.60
Niacinamide	3.02
Pyridoxal HCL	3.00
Pyridoxine HCL	0.031
Riboflavin	0.319
Thiamine HCL	3.17
Thymidine	0.365
Vitamin B ₁₂	0.680

Other Components

HEPES Buffer	25 mM
Na Hypoxanthine	2.39 mg/L
Lipoic Acid	0.105
Sodium Putrescine-2HCL	0.081
Sodium Pyruvate	55.0
Sodium Selenite	0.0067
Ethanolamine	20mM
Ferric Citrate	0.122
Methyl-B-Cyclodextrin complexed with Linoleic Acid	41.70

Methyl-B-Cyclodextrin complexed with Oleic Acid	33.33
Methyl-B-Cyclodextrin complexed with Retinal Acetate	10

Adjust osmolarity to 327 mOsm

Example 12: Construction of GAS Reporter Construct

5 One signal transduction pathway involved in the differentiation and proliferation of cells is called the Jaks-STATs pathway. Activated proteins in the Jaks-STATs pathway bind to gamma activation site "GAS" elements or interferon-sensitive responsive element ("ISRE"), located in the promoter of many genes. The binding of a protein to these elements alter the expression of the associated gene.

10 GAS and ISRE elements are recognized by a class of transcription factors called Signal Transducers and Activators of Transcription, or "STATs." There are six members of the STATs family. Stat1 and Stat3 are present in many cell types, as is Stat2 (as response to IFN-alpha is widespread). Stat4 is more restricted and is not in many cell types though it has been found in T helper class I, cells after treatment with
15 IL-12. Stat5 was originally called mammary growth factor, but has been found at higher concentrations in other cells including myeloid cells. It can be activated in tissue culture cells by many cytokines.

The STATs are activated to translocate from the cytoplasm to the nucleus upon tyrosine phosphorylation by a set of kinases known as the Janus Kinase ("Jaks")
20 family. Jaks represent a distinct family of soluble tyrosine kinases and include Tyk2, Jak1, Jak2, and Jak3. These kinases display significant sequence similarity and are generally catalytically inactive in resting cells.

The Jaks are activated by a wide range of receptors summarized in the Table below. (Adapted from review by Schidler and Darnell, Ann. Rev. Biochem. 64:621-51
25 (1995).) A cytokine receptor family, capable of activating Jaks, is divided into two groups: (a) Class 1 includes receptors for IL-2, IL-3, IL-4, IL-6, IL-7, IL-9, IL-11, IL-12, IL-15, Epo, PRL, GH, G-CSF, GM-CSF, LIF, CNTF, and thrombopoietin; and (b) Class 2 includes IFN-a, IFN-g, and IL-10. The Class 1 receptors share a conserved cysteine motif (a set of four conserved cysteines and one tryptophan) and a
30 WSXWS motif (a membrane proximal region encoding Trp-Ser-Xxx-Trp-Ser (SEQ ID NO:2)).

Thus, on binding of a ligand to a receptor, Jaks are activated, which in turn activate STATs, which then translocate and bind to GAS elements. This entire process is encompassed in the Jaks-STATs signal transduction pathway.

- 5 Therefore, activation of the Jaks-STATs pathway, reflected by the binding of the GAS or the ISRE element, can be used to indicate proteins involved in the proliferation and differentiation of cells. For example, growth factors and cytokines are known to activate the Jaks-STATs pathway. (See Table below.) Thus, by using GAS elements linked to reporter molecules, activators of the Jaks-STATs pathway can be identified.

	<u>Ligand</u>	<u>tyk2</u>	<u>JAKs</u>			<u>STATS</u>	<u>GAS(elements) or ISRE</u>
			<u>Jak1</u>	<u>Jak2</u>	<u>Jak3</u>		
5	<u>IFN family</u>						
	IFN-a/B	+	+	-	-	1,2,3	ISRE
	IFN-g		+	+	-	1	GAS (IRF1>Lys6>IFP)
	Il-10	+	?	?	-	1,3	
10	<u>gp130 family</u>						
	IL-6 (Pleiotrohic)	+	+	+	?	1,3	GAS (IRF1>Lys6>IFP)
	Il-11(Pleiotrohic)	?	+	?	?	1,3	
	OnM(Pleiotrohic)	?	+	+	?	1,3	
	LIF(Pleiotrohic)	?	+	+	?	1,3	
15	CNTF(Pleiotrohic)	-/+	+	+	?	1,3	
	G-CSF(Pleiotrohic)	?	+	?	?	1,3	
	IL-12(Pleiotrohic)	+	-	+	+	1,3	
20	<u>g-C family</u>						
	IL-2 (lymphocytes)	-	+	-	+	1,3,5	GAS
	IL-4 (lymph/myeloid)	-	+	-	+	6	GAS (IRF1 = IFP >>Ly6)(IgH)
	IL-7 (lymphocytes)	-	+	-	+	5	GAS
	IL-9 (lymphocytes)	-	+	-	+	5	GAS
	IL-13 (lymphocyte)	-	+	?	?	6	GAS
	IL-15	?	+	?	+	5	GAS
25	<u>gp140 family</u>						
	IL-3 (myeloid)	-	-	+	-	5	GAS (IRF1>IFP>>Ly6)
	IL-5 (myeloid)	-	-	+	-	5	GAS
30	GM-CSF (myeloid)	-	-	+	-	5	GAS
	<u>Growth hormone family</u>						
	GH	?	-	+	-	5	
35	PRL	?	+/-	+	-	1,3,5	
	EPO	?	-	+	-	5	GAS(B-CAS>IRF1=IFP>>Ly6)
	<u>Receptor Tyrosine Kinases</u>						
40	EGF	?	+	+	-	1,3	GAS (IRF1)
	PDGF	?	+	+	-	1,3	
	CSF-1	?	+	+	-	1,3	GAS (not IRF1)

To construct a synthetic GAS containing promoter element, which is used in the Biological Assays described in Examples 13-14, a PCR based strategy is employed to generate a GAS-SV40 promoter sequence. The 5' primer contains four tandem copies of the GAS binding site found in the IRF1 promoter and previously demonstrated to bind STATs upon induction with a range of cytokines (Rothman et al., Immunity 1:457-468 (1994).), although other GAS or ISRE elements can be used instead. The 5' primer also contains 18bp of sequence complementary to the SV40 early promoter sequence and is flanked with an XhoI site. The sequence of the 5' primer is:

5':GCGCCTCGAGATTTCCCGAAATCTAGATTTCCCGAAATGATTTCCCG
 10 AAATGATTTCCCGAAATATCTGCCATCTCAATTAG:3' (SEQ ID NO:3)

The downstream primer is complementary to the SV40 promoter and is flanked with a Hind III site: 5':GCGGCAAGCTTTTGTCAAAGCCTAGGC:3' (SEQ ID NO:4)

PCR amplification is performed using the SV40 promoter template present in the B-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI/Hind III and subcloned into BLSK2-. (Stratagene.) Sequencing with forward and reverse primers confirms that the insert contains the following sequence:

5':CTCGAGATTTCCCGAAATCTAGATTTCCCGAAATGATTTCCCGAAATG
 20 ATTTCCCGAAATATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCC
 CTAAGTCCGCCCATCCCGCCCCTAAGTCCGCCCAGTTCCGCCCATTCTCCGC
 CCCATGGCTGACTAATTTTTTTTATTTATGCAGAGGCCGAGGCCGCTCGGC
 CTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCCTAGGCTTT
 TGCAAAAAGCTT:3' (SEQ ID NO:5)

With this GAS promoter element linked to the SV40 promoter, a GAS:SEAP2 reporter construct is next engineered. Here, the reporter molecule is a secreted alkaline phosphatase, or "SEAP." Clearly, however, any reporter molecule can be instead of SEAP, in this or in any of the other Examples. Well known reporter molecules that can be used instead of SEAP include chloramphenicol acetyltransferase (CAT), luciferase, alkaline phosphatase, B-galactosidase, green fluorescent protein (GFP), or any protein detectable by an antibody.

The above sequence confirmed synthetic GAS-SV40 promoter element is subcloned into the pSEAP-Promoter vector obtained from Clontech using HindIII and XhoI, effectively replacing the SV40 promoter with the amplified GAS:SV40 promoter element, to create the GAS-SEAP vector. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

Thus, in order to generate mammalian stable cell lines expressing the GAS-SEAP reporter, the GAS-SEAP cassette is removed from the GAS-SEAP vector using SalI and NotI, and inserted into a backbone vector containing the neomycin resistance gene, such as pGFP-1 (Clontech), using these restriction sites in the multiple cloning site, to create the GAS-SEAP/Neo vector. Once this vector is transfected into mammalian cells, this vector can then be used as a reporter molecule for GAS binding as described in Examples 13-14.

Other constructs can be made using the above description and replacing GAS with a different promoter sequence. For example, construction of reporter molecules containing NFK-B and EGR promoter sequences are described in Examples 15 and 16. However, many other promoters can be substituted using the protocols described in these Examples. For instance, SRE, IL-2, NFAT, or Osteocalcin promoters can be substituted, alone or in combination (e.g., GAS/NF-KB/EGR, GAS/NF-KB, IL-2/NFAT, or NF-KB/GAS). Similarly, other cell lines can be used to test reporter construct activity, such as HELA (epithelial), HUVEC (endothelial), Reh (B-cell), Saos-2 (osteoblast), HUVAC (aortic), or Cardiomyocyte.

Example 13: High-Throughput Screening Assay for T-cell Activity.

The following protocol is used to assess T-cell activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate T-cells. T-cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The T-cell used in this assay is Jurkat T-cells (ATCC Accession No. TIB-152), although Molt-3 cells (ATCC Accession No. CRL-1552) and Molt-4 cells (ATCC Accession No. CRL-1582) cells can also be used.

Jurkat T-cells are lymphoblastic CD4+ Th1 helper cells. In order to generate stable cell lines, approximately 2 million Jurkat cells are transfected with the GAS-SEAP/neo vector using DMRIE-C (Life Technologies)(transfection procedure described below). The transfected cells are seeded to a density of approximately 20,000 cells per well and transfectants resistant to 1 mg/ml gentamicin selected. Resistant colonies are expanded and then tested for their response to increasing concentrations of interferon gamma. The dose response of a selected clone is demonstrated.

Specifically, the following protocol will yield sufficient cells for 75 wells containing 200 ul of cells. Thus, it is either scaled up, or performed in multiple to generate sufficient cells for multiple 96 well plates. Jurkat cells are maintained in RPMI + 10% serum with 1%Pen-Strep. Combine 2.5 mls of OPTI-MEM (Life Technologies)

with 10 ug of plasmid DNA in a T25 flask. Add 2.5 ml OPTI-MEM containing 50 ul of DMRIE-C and incubate at room temperature for 15-45 mins.

During the incubation period, count cell concentration, spin down the required number of cells (10^7 per transfection), and resuspend in OPTI-MEM to a final
5 concentration of 10^7 cells/ml. Then add 1ml of 1×10^7 cells in OPTI-MEM to T25 flask and incubate at 37°C for 6 hrs. After the incubation, add 10 ml of RPMI + 15% serum.

The Jurkat:GAS-SEAP stable reporter lines are maintained in RPMI + 10% serum, 1 mg/ml Genticin, and 1% Pen-Strep. These cells are treated with supernatants containing a polypeptide as produced by the protocol described in Example 11.

10 On the day of treatment with the supernatant, the cells should be washed and resuspended in fresh RPMI + 10% serum to a density of 500,000 cells per ml. The exact number of cells required will depend on the number of supernatants being screened. For one 96 well plate, approximately 10 million cells (for 10 plates, 100 million cells) are required.

15 Transfer the cells to a triangular reservoir boat, in order to dispense the cells into a 96 well dish, using a 12 channel pipette. Using a 12 channel pipette, transfer 200 ul of cells into each well (therefore adding 100,000 cells per well).

After all the plates have been seeded, 50 ul of the supernatants are transferred directly from the 96 well plate containing the supernatants into each well using a 12
20 channel pipette. In addition, a dose of exogenous interferon gamma (0.1, 1.0, 10 ng) is added to wells H9, H10, and H11 to serve as additional positive controls for the assay.

The 96 well dishes containing Jurkat cells treated with supernatants are placed in an incubator for 48 hrs (note: this time is variable between 48-72 hrs). 35 ul samples
25 from each well are then transferred to an opaque 96 well plate using a 12 channel pipette. The opaque plates should be covered (using sellophene covers) and stored at -20°C until SEAP assays are performed according to Example 17. The plates containing the remaining treated cells are placed at 4°C and serve as a source of material for repeating the assay on a specific well if desired.

30 As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate Jurkat T cells. Over 30 fold induction is typically observed in the positive control wells.

Example 14: High-Throughput Screening Assay Identifying Myeloid Activity

The following protocol is used to assess myeloid activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate myeloid cells.

- 5 Myeloid cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The myeloid cell used in this assay is U937, a pre-monocyte cell line, although TF-1, HL60, or KG1 can be used.

- 10 To transiently transfect U937 cells with the GAS/SEAP/Neo construct produced in Example 12, a DEAE-Dextran method (Kharbanda et. al., 1994, Cell Growth & Differentiation, 5:259-265) is used. First, harvest 2×10^7 U937 cells and wash with PBS. The U937 cells are usually grown in RPMI 1640 medium containing 10% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 mg/ml streptomycin.

- 15 Next, suspend the cells in 1 ml of 20 mM Tris-HCl (pH 7.4) buffer containing 0.5 mg/ml DEAE-Dextran, 8 ug GAS-SEAP2 plasmid DNA, 140 mM NaCl, 5 mM KCl, 375 uM $\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$, 1 mM MgCl_2 , and 675 uM CaCl_2 . Incubate at 37°C for 45 min.

- 20 Wash the cells with RPMI 1640 medium containing 10% FBS and then resuspend in 10 ml complete medium and incubate at 37°C for 36 hr.

The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 400 ug/ml G418 for couple of passages.

- 25 These cells are tested by harvesting 1×10^8 cells (this is enough for ten 96-well plates assay) and wash with PBS. Suspend the cells in 200 ml above described growth medium, with a final density of 5×10^5 cells/ml. Plate 200 ul cells per well in the 96-well plate (or 1×10^5 cells/well).

- 30 Add 50 ul of the supernatant prepared by the protocol described in Example 11. Incubate at 37°C for 48 to 72 hr. As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate U937 cells. Over 30 fold induction is typically observed in the positive control wells. SEAP assay the supernatant according to the protocol described in Example 17.

- 35 **Example 15: High-Throughput Screening Assay Identifying Neuronal Activity.**

When cells undergo differentiation and proliferation, a group of genes are activated through many different signal transduction pathways. One of these genes, EGR1 (early growth response gene 1), is induced in various tissues and cell types upon activation. The promoter of EGR1 is responsible for such induction. Using the EGR1
5 promoter linked to reporter molecules, activation of cells can be assessed.

Particularly, the following protocol is used to assess neuronal activity in PC12 cell lines. PC12 cells (rat phenochromocytoma cells) are known to proliferate and/or differentiate by activation with a number of mitogens, such as TPA (tetradecanoyl phorbol acetate), NGF (nerve growth factor), and EGF (epidermal growth factor). The
10 EGR1 gene expression is activated during this treatment. Thus, by stably transfecting PC12 cells with a construct containing an EGR promoter linked to SEAP reporter, activation of PC12 cells can be assessed.

The EGR/SEAP reporter construct can be assembled by the following protocol. The EGR-1 promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene 6:867-871
15 (1991)) can be PCR amplified from human genomic DNA using the following primers:

5' GCGCTCGAGGGATGACAGCGATAGAACCCCGG -3' (SEQ ID NO:6)

5' GCGAAGCTTCGCGACTCCCCGGATCCGCCTC-3' (SEQ ID NO:7)

Using the GAS:SEAP/Neo vector produced in Example 12, EGR1 amplified product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector
20 using restriction enzymes XhoI/HindIII, removing the GAS/SV40 stuffer. Restrict the EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1 promoter.

To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30 dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter
25 sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and allowed to air dry for 2 hr.

PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker) containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100
30 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is done every three to four days. Cells are removed from the plates by scraping and resuspended with pipetting up and down for more than 15 times.

Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine protocol described in Example 11. EGR-SEAP/PC12 stable cells are obtained by
35 growing the cells in 300 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 300 ug/ml G418 for couple of passages.

To assay for neuronal activity, a 10 cm plate with cells around 70 to 80% confluent is screened by removing the old medium. Wash the cells once with PBS (Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

- 5 The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as 5×10^5 cells/ml.

- 10 Add 200 ul of the cell suspension to each well of 96-well plate (equivalent to 1×10^5 cells/well). Add 50 ul supernatant produced by Example 11, 37°C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ul of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 17.

15

Example 16: High-Throughput Screening Assay for T-cell Activity

- NF- κ B (Nuclear Factor κ B) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by
20 expression of certain viral gene products. As a transcription factor, NF- κ B regulates the expression of genes involved in immune cell activation, control of apoptosis (NF- κ B appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

- 25 In non-stimulated conditions, NF- κ B is retained in the cytoplasm with I- κ B (Inhibitor κ B). However, upon stimulation, I- κ B is phosphorylated and degraded, causing NF- κ B to shuttle to the nucleus, thereby activating transcription of target genes. Target genes activated by NF- κ B include IL-2, IL-6, GM-CSF, ICAM-1 and class 1 MHC.

- 30 Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF- κ B promoter element are used to screen the supernatants produced in Example 11. Activators or inhibitors of NF- κ B would be useful in treating

diseases. For example, inhibitors of NF- κ B could be used to treat those diseases related to the acute or chronic activation of NF- κ B, such as rheumatoid arthritis.

To construct a vector containing the NF- κ B promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF- κ B binding site (GGGGACTTTCCC) (SEQ ID NO:8), 18 bp of sequence complementary to the 5' end of the SV40 early promoter sequence, and is flanked with an XhoI site:
 5':GCGGCCTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGAC
 TTTCCATCCTGCCATCTCAATTAG:3' (SEQ ID NO:9)

The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:
 5':GCGGCAAGCTTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO:4)

PCR amplification is performed using the SV40 promoter template present in the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2-. (Stratagene)
 15 Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

5':CTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGGACTTTCC
 ATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCCCCTAACTCCGCCCCA
 20 TCCCGCCCCCTAACTCCGCCCCAGTTCCGCCCCATTCTCCGCCCCATGGCTGACT
 AATTTTTTTTTATTTATGCAGAGGCCGAGGCCGCTCGGCCTCTGAGCTATTC
 CAGAAGTAGTGAGGAGGCTTTTTTTGGAGGCCTAGGCTTTTGCAAAAAGCTT:
 3' (SEQ ID NO:10)

25 Next, replace the SV40 minimal promoter element present in the pSEAP2- promoter plasmid (Clontech) with this NF- κ B/SV40 fragment using XhoI and HindIII. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

In order to generate stable mammalian cell lines, the NF- κ B/SV40/SEAP
 30 cassette is removed from the above NF- κ B/SEAP vector using restriction enzymes SalI and NotI, and inserted into a vector containing neomycin resistance. Particularly, the NF- κ B/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP gene, after restricting pGFP-1 with SalI and NotI.

Once NF- κ B/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are created and maintained according to the protocol described in Example 13. Similarly, the method for assaying supernatants with these stable Jurkat T-cells is also described in Example 13. As a positive control, exogenous TNF alpha (0.1, 1, 10 ng) is added to wells H9, H10, and H11, with a 5-10 fold activation typically observed.

Example 17: Assay for SEAP Activity

As a reporter molecule for the assays described in Examples 13-16, SEAP activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the following general procedure. The Tropix Phospho-light Kit supplies the Dilution, Assay, and Reaction Buffers used below.

Prime a dispenser with the 2.5x Dilution Buffer and dispense 15 μ l of 2.5x dilution buffer into Optiplates containing 35 μ l of a supernatant. Seal the plates with a plastic sealer and incubate at 65°C for 30 min. Separate the Optiplates to avoid uneven heating.

Cool the samples to room temperature for 15 minutes. Empty the dispenser and prime with the Assay Buffer. Add 50 μ l Assay Buffer and incubate at room temperature 5 min. Empty the dispenser and prime with the Reaction Buffer (see the table below). Add 50 μ l Reaction Buffer and incubate at room temperature for 20 minutes. Since the intensity of the chemiluminescent signal is time dependent, and it takes about 10 minutes to read 5 plates on luminometer, one should treat 5 plates at each time and start the second set 10 minutes later.

Read the relative light unit in the luminometer. Set H12 as blank, and print the results. An increase in chemiluminescence indicates reporter activity.

Reaction Buffer Formulation:

# of plates	Rxn buffer diluent (ml)	CSPD (ml)
10	60	3
11	65	3.25
12	70	3.5
13	75	3.75
14	80	4
15	85	4.25
16	90	4.5
17	95	4.75
18	100	5
19	105	5.25
20	110	5.5
21	115	5.75
22	120	6

23	125	6.25
24	130	6.5
25	135	6.75
26	140	7
27	145	7.25
28	150	7.5
29	155	7.75
30	160	8
31	165	8.25
32	170	8.5
33	175	8.75
34	180	9
35	185	9.25
36	190	9.5
37	195	9.75
38	200	10
39	205	10.25
40	210	10.5
41	215	10.75
42	220	11
43	225	11.25
44	230	11.5
45	235	11.75
46	240	12
47	245	12.25
48	250	12.5
49	255	12.75
50	260	13

Example 18: High-Throughput Screening Assay Identifying Changes in Small Molecule Concentration and Membrane Permeability

Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane potential. These alterations can be measured in an assay to identify supernatants which bind to receptors of a particular cell. Although the following protocol describes an assay for calcium, this protocol can easily be modified to detect changes in potassium, sodium, pH, membrane potential, or any other small molecule which is detectable by a fluorescent probe.

The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small molecules. Clearly, any fluorescent molecule detecting a small molecule can be used instead of the calcium fluorescent molecule, fluo-3, used here.

For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black 96-well plate with clear bottom. The plate is incubated in a CO₂ incubator for 20 hours. The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash.

A stock solution of 1 mg/ml fluo-3 is made in 10% pluronic acid DMSO. To load the cells with fluo-3, 50 μ l of 12 μ g/ml fluo-3 is added to each well. The plate is incubated at 37°C in a CO₂ incubator for 60 min. The plate is washed four times in the Biotek washer with HBSS leaving 100 μ l of buffer.

- 5 For non-adherent cells, the cells are spun down from culture media. Cells are re-suspended to 2-5x10⁶ cells/ml with HBSS in a 50-ml conical tube. 4 μ l of 1 mg/ml fluo-3 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. The tube is then placed in a 37°C water bath for 30-60 min. The cells are washed twice with HBSS, resuspended to 1x10⁶ cells/ml, and dispensed into a microplate, 100
10 μ l/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley CellWash with 200 μ l, followed by an aspiration step to 100 μ l final volume.

For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-3. The supernatant is added to the well, and a change in fluorescence is detected.

- To measure the fluorescence of intracellular calcium, the FLIPR is set for the
15 following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and (6) Sample addition is 50 μ l. Increased emission at 530 nm indicates an extracellular signaling event which has resulted in an increase in the intracellular Ca⁺⁺ concentration.

20

Example 19: High-Throughput Screening Assay Identifying Tyrosine Kinase Activity

- The Protein Tyrosine Kinases (PTK) represent a diverse group of transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase
25 RPTK) group are receptors for a range of mitogenic and metabolic growth factors including the PDGF, FGF, EGF, NGF, HGF and Insulin receptor subfamilies. In addition there are a large family of RPTKs for which the corresponding ligand is unknown. Ligands for RPTKs include mainly secreted small proteins, but also membrane-bound and extracellular matrix proteins.

- 30 Activation of RPTK by ligands involves ligand-mediated receptor dimerization, resulting in transphosphorylation of the receptor subunits and activation of the cytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and non-receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members
35 of which mediate signal transduction triggered by the cytokine superfamily of receptors (e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

Because of the wide range of known factors capable of stimulating tyrosine kinase activity, the identification of novel human secreted proteins capable of activating tyrosine kinase signal transduction pathways are of interest. Therefore, the following protocol is designed to identify those novel human secreted proteins capable of
5 activating the tyrosine kinase signal transduction pathways.

Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyne Silent Screen Plates purchased from Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 100% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr
10 with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or 10% Matrigel purchased from Becton Dickinson (Bedford, MA), or calf serum, rinsed with PBS and stored at 4°C. Cell growth on these plates is assayed by seeding 5,000 cells/well in growth medium and indirect quantitation of cell number through use of
15 alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford, MA) are used to cover the Loprodyne Silent Screen Plates. Falcon Microtest III cell culture plates can also be used in some proliferation experiments.

To prepare extracts, A431 cells are seeded onto the nylon membranes of
20 Loprodyne plates (20,000/200ml/well) and cultured overnight in complete medium. Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example 11, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na₃VO₄, 2 mM Na₄P₂O₇
25 and a cocktail of protease inhibitors (# 1836170) obtained from Boehringer Mannheim (Indianapolis, IN) is added to each well and the plate is shaken on a rotating shaker for 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract filtered through the 0.45 mm membrane bottoms of each well using house vacuum. Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum
30 manifold and immediately placed on ice. To obtain extracts clarified by centrifugation, the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4°C at 16,000 x g.

Test the filtered extracts for levels of tyrosine kinase activity. Although many methods of detecting tyrosine kinase activity are known, one method is described here.

35 Generally, the tyrosine kinase activity of a supernatant is evaluated by determining its ability to phosphorylate a tyrosine residue on a specific substrate (a

biotinylated peptide). Biotinylated peptides that can be used for this purpose include PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and PS1 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for a range of tyrosine kinases and are available from Boehringer Mannheim.

- 5 The tyrosine kinase reaction is set up by adding the following components in order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg₂⁺ (5mM ATP/50mM MgCl₂), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride, pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl₂, 5 mM MnCl₂, 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the
- 10 components gently and preincubate the reaction mix at 30°C for 2 min. Initiate the reaction by adding 10ul of the control enzyme or the filtered supernatant.

The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mM EDTA and place the reactions on ice.

- Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction
- 15 mixture to a microtiter plate (MTP) module and incubating at 37°C for 20 min. This allows the streptavidin coated 96 well plate to associate with the biotinylated peptide. Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of anti-phosphotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-POD(0.5u/ml)) to each well and incubate at 37°C for one hour. Wash the well as
- 20 above.

- Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and incubate at room temperature for at least 5 mins (up to 30 min). Measure the absorbance of the sample at 405 nm by using ELISA reader. The level of bound peroxidase activity is quantitated using an ELISA reader and reflects the level of
- 25 tyrosine kinase activity.

Example 20: High-Throughput Screening Assay Identifying Phosphorylation Activity

- As a potential alternative and/or complement to the assay of protein tyrosine
- 30 kinase activity described in Example 19, an assay which detects activation (phosphorylation) of major intracellular signal transduction intermediates can also be used. For example, as described below one particular assay can detect tyrosine phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase,
- 35 Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other

phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by substituting these molecules for Erk-1 or Erk-2 in the following assay.

Specifically, assay plates are made by coating the wells of a 96-well ELISA plate with 0.1ml of protein G (1ug/ml) for 2 hr at room temp, (RT). The plates are then
5 rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates are then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1 and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this step can easily be modified by substituting a monoclonal antibody detecting any of the above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4°C
10 until use.

A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and cultured overnight in growth medium. The cells are then starved for 48 hr in basal medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants obtained in Example 11 for 5-20 minutes. The cells are then solubilized and extracts
15 filtered directly into the assay plate.

After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a positive control, a commercial preparation of MAP kinase (10ng/well) is used in place of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody (1ug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and
20 Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The bound polyclonal antibody is then quantitated by successive incubations with Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over background indicates a phosphorylation.

25

Example 21: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is be isolated. cDNA is then generated from
30 these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in SEQ ID NO:X. Suggested PCR conditions consist of 35 cycles at 95°C for 30 seconds; 60-120 seconds at 52-58°C; and 60-120 seconds at 70°C, using buffer solutions described in Sidransky, D., et al., Science 252:706 (1991).

35 PCR products is then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase. (Epicentre Technologies).

The intron-exon borders of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations is then cloned and sequenced to validate the results of the direct sequencing.

5 PCR products is cloned into T-tailed vectors as described in Holton, T.A. and Graham, M.W., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals is identified by mutations not present in unaffected individuals.

10 Genomic rearrangements are also observed as a method of determining alterations in a gene corresponding to a polynucleotide. Genomic clones isolated according to Example 2 are nick-translated with digoxigenindeoxy-uridine 5'-triphosphate (Boehringer Mannheim), and FISH performed as described in Johnson, Cg. et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the corresponding genomic locus.

15 Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson, Cv. et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and
20 chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosome alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and translocations. These alterations are used as a diagnostic marker for an associated
25 disease.

Example 22: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

30 A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

35 For example, antibody-sandwich ELISAs are used to detect soluble polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method

described in Example 10. The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, serial dilutions of the sample should be used to
5 validate results. The plates are then washed three times with deionized or distilled water to remove unbounded polypeptide.

Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove
10 unbounded conjugate.

Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on
15 the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale). Interpolate the concentration of the polypeptide in the sample using the standard curve.

Example 23: Formulating a Polypeptide

The secreted polypeptide composition will be formulated and dosed in a fashion
20 consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with the secreted polypeptide alone), the site of delivery, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

As a general proposition, the total pharmaceutically effective amount of secreted polypeptide administered parenterally per dose will be in the range of about 1 µg/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If
30 given continuously, the secreted polypeptide is typically administered at a dose rate of about 1 µg/kg/hour to about 50 µg/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending
35 on the desired effect.

Pharmaceutical compositions containing the secreted protein of the invention are administered orally, rectally, parenterally, intracisternally, intravaginally,

intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), buccally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

The secreted polypeptide is also suitably administered by sustained-release systems. Suitable examples of sustained-release compositions include semi-permeable polymer matrices in the form of shaped articles, e.g., films, or microcapsules. Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman, U. et al., Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (R. Langer et al., J. Biomed. Mater. Res. 15:167-277 (1981), and R. Langer, Chem. Tech. 12:98-105 (1982)), ethylene vinyl acetate (R. Langer et al.) or poly-D- (-)-3-hydroxybutyric acid (EP 133,988). Sustained-release compositions also include liposomally entrapped polypeptides. Liposomes containing the secreted polypeptide are prepared by methods known per se: DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. USA 82:3688-3692 (1985); Hwang et al., Proc. Natl. Acad. Sci. USA 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal secreted polypeptide therapy.

For parenteral administration, in one embodiment, the secreted polypeptide is formulated generally by mixing it at the desired degree of purity, in a unit dosage injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations employed and is compatible with other ingredients of the formulation. For example, the formulation preferably does not include oxidizing agents and other compounds that are known to be deleterious to polypeptides.

Generally, the formulations are prepared by contacting the polypeptide uniformly and intimately with liquid carriers or finely divided solid carriers or both. Then, if necessary, the product is shaped into the desired formulation. Preferably the carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood of the recipient. Examples of such carrier vehicles include water, saline, Ringer's solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl oleate are also useful herein, as well as liposomes.

The carrier suitably contains minor amounts of additives such as substances that enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at the dosages and concentrations employed, and include buffers such as phosphate, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, manose, or dextrans; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates, poloxamers, or PEG.

The secreted polypeptide is typically formulated in such vehicles at a concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.

Any polypeptide to be used for therapeutic administration can be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutic polypeptide compositions generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

Polypeptides ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous polypeptide solution, and the resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the lyophilized polypeptide using bacteriostatic Water-for-Injection.

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the present invention may be employed in conjunction with other therapeutic compounds.

Example 24: Method of Treating Decreased Levels of the Polypeptide

It will be appreciated that conditions caused by a decrease in the standard or normal expression level of a secreted protein in an individual can be treated by administering the polypeptide of the present invention, preferably in the secreted form. Thus, the invention also provides a method of treatment of an individual in need of an increased level of the polypeptide comprising administering to such an individual a pharmaceutical composition comprising an amount of the polypeptide to increase the activity level of the polypeptide in such an individual.

For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the polypeptide for six consecutive days. Preferably, the polypeptide is in the secreted form. The exact details of the dosing scheme, based on administration and formulation, are provided in Example 23.

Example 25: Method of Treating Increased Levels of the Polypeptide

Antisense technology is used to inhibit production of a polypeptide of the present invention. This technology is one example of a method of decreasing levels of a polypeptide, preferably a secreted form, due to a variety of etiologies, such as cancer.

For example, a patient diagnosed with abnormally increased levels of a polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, 2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is provided in Example 23.

Example 26: Method of Treatment Using Gene Therapy

One method of gene therapy transplants fibroblasts, which are capable of expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and separated into small pieces. Small chunks of the tissue are placed on a wet surface of a tissue culture flask, approximately ten pieces are placed in each flask. The flask is turned upside down, closed tight and left at room temperature over night. After 24 hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to the bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS, penicillin and streptomycin, is added. The flasks are then incubated at 37°C for approximately one week.

At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long
5 terminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is fractionated on agarose gel and purified, using glass beads.

The cDNA encoding a polypeptide of the present invention can be amplified using PCR primers which correspond to the 5' and 3' end sequences respectively as set
10 forth in Example 1. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to
15 transform bacteria HB101, which are then plated onto agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is
20 then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media,
25 containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is
30 required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is being produced.

The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

35

It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

- 5 The entire disclosure of each document cited (including patents, patent applications, journal articles, abstracts, laboratory manuals, books, or other disclosures) in the Background of the Invention, Detailed Description, and Examples is hereby incorporated herein by reference.

(1) GENERAL INFORMATION:

(i) APPLICANTS: Human Genome Sciences, Inc. et al.

(ii) TITLE OF INVENTION: 70 Human Secreted Proteins

5 (iii) NUMBER OF SEQUENCES: 273

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(v) COMPUTER READABLE FORM:

15 (A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage

(B) COMPUTER: HP Vectra 486/33

(C) OPERATING SYSTEM: MSDOS version 6.2

(D) SOFTWARE: ASCII Text

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160

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5 (2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 733 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

10 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

	GGGATCCGGA GCCCAAATCT TCTGACAAAA CTCACACATG CCCACCGTGC CCAGCACCTG	60
	AATTCGAGGG TGCACCGTCA GTCTTCCTCT TCCCCCAAA ACCCAAGGAC ACCCTCATGA	120
15	TCTCCCGGAC TCCTGAGGTC ACATGCGTGG TGGTGGACGT AAGCCACGAA GACCCTGAGG	180
	TCAAGTTCAA CTGGTACGTG GACGGCGTGG AGGTGCATAA TGCCAAGACA AAGCCGCGGG	240
	AGGAGCAGTA CAACAGCACG TACCGTGTGG TCAGCGTCCT CACCGTCCTG CACCAGGACT	300
	GGCTGAATGG CAAGGAGTAC AAGTGCAAGG TCTCCAACAA AGCCCTCCCA ACCCCCATCG	360
	AGAAAACCAT CTCCAAAGCC AAAGGGCAGC CCCGAGAACC ACAGGTGTAC ACCCTGCCCC	420
20	CATCCCGGGA TGAGCTGACC AAGAACCAGG TCAGCCTGAC CTGCCTGGTC AAAGGCTTCT	480
	ATCCAAGCGA CATCGCCGTG GACTGGGAGA GCAATGGGCA GCCGGAGAAC AACTACAAGA	540
	CCACGCCTCC CGTGCTGGAC TCCGACGGCT CTTCTTCCT CTACAGCAAG CTCACCGTGG	600
	ACAAGAGCAG GTGGCAGCAG GGAACGTCT TCTCATGCTC CGTGATGCAT GAGGCTCTGC	660
	ACAACCACTA CACGCAGAAG AGCCTCTCCC TGTCTCCGGG TAAATGAGTG CGACGGCCGC	720
25	GACTCTAGAG GAT	733

(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 5 amino acids

30 (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

Trp Ser Xaa Trp Ser

5 1 5

(2) INFORMATION FOR SEQ ID NO: 3:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 86 base pairs

10 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

15 GCGCCTCGAG ATTTCCCCGA AATCTAGATT TCCCCGAAAT GATTCCCCG AAATGATTTC 60
CCCGAAATAT CTGCCATCTC AATTAG 86

(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:

20 (A) LENGTH: 27 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

25 GCGGCAAGCT TTTTGCAAAG CCTAGGC 27

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 271 base pairs

30 (B) TYPE: nucleic acid

162

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

CTCGAGATTT CCCCAGAAATC TAGATTTCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG 60
5 AAATATCTGC CATCTCAATT AGTCAGCAAC CATAGTCCCG CCCCTAACTC CGCCCATCCC 120
GCCCCTAACT CCGCCAGTT CCGCCATTC TCCGCCCAT GGCTGACTAA TTTTTTTTAT 180
TTATGCAGAG GCCGAGGCCG CCTCGGCCTC TGAGCTATTC CAGAAGTAGT GAGGAGGCTT 240
TTTTGGAGGC CTAGGCTTTT GCAAAAAGCT T 271

10 (2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 32 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

15 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

GCGCTCGAGG GATGACAGCG ATAGAACCCC GG 32

20 (2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 31 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

25 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

GCGAAGCTTC GCGACTCCCC GGATCCGCCT C 31

30 (2) INFORMATION FOR SEQ ID NO: 8:

163

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 12 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

5 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

GGGGACTTTC CC

12

10 (2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 73 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

15 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

GCGGCCTCGA GGGGACTTTC CCGGGGACTT TCCGGGGACT TTCCGGGACT TTCCATCCTG

60

CCATCTCAAT TAG

73

20

(2) INFORMATION FOR SEQ ID NO: 10:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 256 base pairs

25 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

30 CTCGAGGGGA CTTTCCCGG GACTTTCCGG GGACTTTCCG GAGACTTTCCA TCTGCCATCT

60

164

CAATTAGTCA GCAACCATAG TCCCGCCCCCT AACTCCGCCC ATCCCGCCCC TAACTCCGCC 120
 CAGTTCCGCC CATTTCTCCGC CCCATGGCTG ACTAATTTTT TTTATTTATG CAGAGGCCGA 180
 GGCCGCCTCG GCCTCTGAGC TATTCCAGAA GTAGTGAGGA GGCTTTTTTTG GAGGCCTAGG 240
 CTTTGTGCAA AAGCTT 256

5

(2) INFORMATION FOR SEQ ID NO: 11:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1739 base pairs

(B) TYPE: nucleic acid

10

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

GGCCTCCCGA GGCCGCGGGA CCTGCAGAGA GGACAGCCGG CCTGCGCCGG GACATGCGGC 60
 15 CCCAGGAGCT CCCCAGGCTC GCGTTCCCGT TGCTGTGTTT GCTGTTGCTG CTGCTGCCGC 120
 CGCCGCCGTG CCCTGCCAC AGCGCCACGC GTTTCGACCC CACCTGGGAG TCCCTGGACG 180
 CCCGCCAGCT GCCCGCGTGG TTTGACCAGG CCAAGTTCGG CATCTTCATC CACTGGGGAG 240
 TGTTTTCCGT GCCCAGCTTC GGTAGCGAGT GGTTCGTGGT GTATTGGCAA AAGGAAAAGA 300
 TACCGAAGTA TGTGGAATTT ATGAAAGATA ATTACCCTCC TARTTTCAAA TATGAAGATT 360
 20 TTGGACCACT ATTTACAGCA AAATTTTTTA ATGCCAACCA RTGGGCARAT ATTTTTCAGG 420
 CCTCTGGTGC CAAATACATT GTCTTAACTT CCAAACATCA TGAAGGCTTT ACCTTGTTGG 480
 GGTCAGAATA TTCGTGGAAC TGAATGCCA TAGATGAGGG GCCCAAGAGG GACATTGTCA 540
 AGGAACTTGA GGTAGCCATT AGGAACAGAA CTGACCTGCG TTTTGGACTG TACTATTCCT 600
 TTTTGAATG GTTTCATCCG CTCTTCCTTG AGGATGAATC CAGTTCATTC CATAAGCGGC 660
 25 AATTTCCAGT TTCTAAGACA TTGCCAGAGC TCTATGAGTT AGTGAACAAC TATCAGCCTG 720
 AGGTTCGTG GTCGGATGGT GACGGAGGAG CACCGGATCA ATACTGGAAC ANCACAGGCT 780
 TCTTGGCCTG GTTATATAAT GAAAGCCAG TTCGGGGCAC AGTAGTCACC AATGATCGTT 840
 GGGGAGCTGG TAGCATCTGT AAGCATGGTG GCTTCTATAC CTCTAGTGTAT CTTTATAACC 900
 CAGGACATCT TTTGCCACAT AAATGGGAAA ACTGCATGAG AATAGACAAA CTGTCTCTGG 960
 30 GCTATAGGAG GGAAGCTGGA ATCTCTGACT ATCTTACAAT TGAAGAATTG GTGAAGCAAC 1020

165

	TTGTAGAGAC AGTTTCATGT GGAGGAAATC TTTTGATGAA TATTGGGCCC AACTAGATG	1080
	GCACCATTC TGTAGTTTTT GAGGAGCGAC TGAGGCAAAT GGGGTCCTGG CTAAAAGTCA	1140
	ATGGAGAAGC TATTTATGAA ACCCATACCT GGCGATCCCA GAATGACACT GTCACCCAG	1200
	ATGTGTGGTA CACATCCAAG CCTAAAGAAA AATTAGTCTA TGCCATTTTT CTAAATGGC	1260
5	CCACATCAGG ACAGCTGTTC CTTGGCCATC CCAAAGCTAT TCTGGGGCA ACAGAGGTGA	1320
	AACTACTGGG CCATGGACAG CCACTTAACT GGATTTCTTT GGAGCAAAAT GGCATTATGG	1380
	TAGAACTGCC ACAGCTAACC ATTCATCAGA TGCCGTGTAA ATGGGGCTGG GCTCTAGCCC	1440
	TRACTAATGT GATCTAAAGT GCAGCAGAGT GGCTGATGCT GCAAGTTATG TCTAAGGCTA	1500
	GGAECTATCA GGTGTCTATA ATTGTAGCAC ATGGAGAAAG CAAATGTAA ACTGGATAAG	1560
10	AAAATTATTT TGGCAGTTCA GCCCTTTCCC TTTTCCCAC TAAATTTTTT CTAAATTAC	1620
	CCATGTAACC ATTTTAACTC TCCAGTGCAC TTGCCATTA AAGTCTCTC ACATTGAAAA	1680
	AAAAAAAAA AAAAACCCCG GGGGGGGGGC CCGGNACCC CATTTGCCCC NTAAAGGGG	1739

(2) INFORMATION FOR SEQ ID NO: 12:

15

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 844 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

	GGCCCCCTGGG CCCGAGGGGC TGGAGCCGGG CCGGGCGGAT GTGGAGCGCG GGCCGCGGCG	60
	GGGCTGCCTG GCCGGTGCTG TTGGGGCTGC TGCTGGCGCT GTTAGTGCCG GGCGGTGGTG	120
	CCGCCAAGAC CGGTGCGGAG CTCGTGACCT GCGGGTCGGT GCTGAAGCTG CTCAATACGC	180
	ACCACCGCGT GCGGCTGCAC TCGCACGACA TCAAATACCG ATTGGGCAGC GGCAGCAAT	240
25	CGGTGACCGG CGTAGAGGCG TCGGACGACG CCAATAGCTA CTGGGGGATC CGCGGCGGCT	300
	CGGAGGGCGG GTGCCGCCG GGGTCCCCG TGCGCTGCTG GTTAGGGGTG AGGCTCACGC	360
	ATGTGCTTAC GGGCAAGAAC CTGCACACGC ACCACTTCTG GTTAGGGGTG TCCAAACAACC	420
	AGGAGGTGAG TGCCTTTGGG GAAGACGGCG AGGGCGAGCT GTTAGGGGTG TCGACAGTGC	480
	GCTGCTCTGG ACAGCACTGG GAGCGTGAGG CTGCTGTGCT GTTAGGGGTG GTGGGCACCT	540
30	CTGTGTTCTT GTCAGTCACG GGTGAGCAGT ATGGAAGCTG GTTAGGGGTG CAGCATGAGG	600

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TCCACGGCAT GCCCAGTGCC AACACGCACA ATACGTGGAA GGCCATGGAA GGCATCTTCA 660
 TCAAGCCTAG TGTGGAGCCC TCTGCAGGTC ACGATGAACT CTGAGTGTGT GGATGGATGG 720
 GTGGATGGAG GGTGGCAGGT GGGGCGTCTG CAGGGCCACT CTTGGCAGAG ACTTTGGGTT 780
 TGTAGGGGTC CTCAAGTGCC TTTGTGATTA AAGAATGTTG GTCTATGAAA AAAAAAAAAA 840
 5 AAAA 844

(2) INFORMATION FOR SEQ ID NO: 13:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 776 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:

TTCGAAATAA AAGATCTGCT CAAGAGAGCC GCAGAAAAAG AAGGTGTATG TTGGGGGTTT 60
 15 AGAGAGCAGG GTCTTGAAAT ACACAGCCCA GAATATGGAG CTTCAGAACA AAGTACAGCT 120
 TCTGGAGGAA CAGAATTGTG CCCTTCTAGA TCAACTGAGG AACTCCAGG CCATGGTGAT 180
 TGAGATATCA AACAAAACCA GCAGCAGCAG CACCTGCATC TTGGTCCTAC TAGTCTCCTT 240
 CTGCCTCCTC CTTGTACCTG CTATGTACTC CTCTGACACA AGGGGGAGCC TGCCAGCTGA 300
 GCATGGAGTG TTGTCCCGCC AGCTTCGTGC CCTCCCCAGT GAGGACCCCTT ACCAGCTGGA 360
 20 GCTGCCTGCC CTGCAGTCAG AAGTGCCGAA AGACAGCACA CACCAGTGGT TGGACGGCTC 420
 AGACTGTGTA CTCCAGGCCC CTGGCAACAC TTCCTGCCTG CTGCATTACA TGCCTCAGGC 480
 TCCCAGTGCA GAGCCTCCCC TGGAGTGGCC ATTCCCTGAC CTCTTCTCAG AGCCTCTCTG 540
 CCGAGGTCCC ATCCTCCCCC TGCAGGCAAA TCTCACAAGG AAGGAGGAT GGCTTCCTAC 600
 25 TGGTAGCCCC TCTGTCATTT TGCAGGACAG ATACTCAGGC TACATATGAG GATATGTGGG 660
 GGGTCTCAGC AGGAGCCTGG GGGGCTCCCC ATCTGTGTCT AAATAAAAAG CGGTGGGCAA 720
 GGGCTGGCCG CAGCTCCTGT GCCCTGTCAG GACGACTGAG ATTCAAAACA CACCAC 776

(2) INFORMATION FOR SEQ ID NO: 14:

- (i) SEQUENCE CHARACTERISTICS:

167

(A) LENGTH: 1376 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

	GAATTCGGCA CGAGGCGCCT ACCCTGCCTG CAGGTGAGCA GTGGTGTGTG AGAGCCAGGC	60
	GTCCCTCTGC CTGCCCCTC AGTGGCAACA CCCGGGAGCT GTTTTGTCTT TTGTGGAGCC	120
	TCAGCAGTTC CCTCTTTCAG AACTCACTGC CAAGAGCCCT GAACAGGAGC CACCATGCAG	180
	TGCTTCAGCT TCATTAAGAC CATGATGATC CTCTCAATT TGCTCATCTT TCTGTGTGGT	240
10	GCAGCCCTGT TGGCAGTGGG CATCTGGGTG TCAATCGATG GGGCATCCTT TCTGAAGATC	300
	TTCTGGGCCAC TGTCGTCCAG TGCCATGCAG TTTGTCAACG TGGGCTACTT CCTCATCGCA	360
	GCCGGCGTTG TGGTCTTTGC TCTTGGTTTC CTGGGCTGCT ATGGTGCTAA GACTGAGAGC	420
	AAGTGTGCCC TCGTGACGTT CTTCTTCATC CTCCTCCTCA TCTTCATGTC TGAGGTTGCA	480
	GCTGCTGTGG TCGCCTTGGT GTACACCACA ATGGCTGAGC ACTTCCTGAC GTTGCTGGTA	540
15	GTGCCTGCCA TCAAGAAAGA TTATGGTTCC CAGGAAGACT TCACTCAAGT GTGGAACACC	600
	ACCATGAAAG GGCTCAAGTG CTGTGGCTTC ACCAACTATA CGGATTTTGA GGACTCACCC	660
	TACTTCAAAG AGAACAGTGC CTTTCCCCCA TTCTGTTGCA ATGACAACGT CACCAACACA	720
	GCCAATGAAA CCTGCACCAA GCAAAAGGCT CACGACCAAA AAGTAGAGGG TTGCTTCAAT	780
	CAGCTTTTGT ATGACATCCG AACTAATGCA GTCACCGTGG GTGGTGTGGC AGCTGGAATT	840
20	GGGGGCCTCG AGCTGGCTGC CATGATTGTG TCCATGTATC TGTACTGCAA TCTACAATAA	900
	GTCCACTTCT GCCTCTGCCA CTA CTGCTGC CACATGGGAA CTGTGAAGAG GCACCCTGGC	960
	AAGCAGCAGT GATTGGGGGA GGGGACAGGA TCTAACAATG TCACTTGGGC CAGAATGGAC	1020
	CTGCCCTTTC TGCTCCAGAC TTGGGGCTAG ATAGGGACCA CTCCTTTTAN GCGATGCCTG	1080
	ACTTTCCTTC CATTGGTGGG TGGATGGGTG GGGGSCATTC CAGAGCCTCT AAGGTAGCCA	1140
25	GTTCTGTTGC CCATTCCCCC AGTCTATTAA ACCCTTGATA TGCCCCCTAG GCCTAGTGGT	1200
	GATCCCAGTG CTCTACTGGG GGATGAGAGA AAGGCATTTT ATAGCCTGGG CATAAGTGAA	1260
	ATCAGCAGAG CCTCTGGGTG GATGTGTAGA AGGCACATCA AAATGCATAA ACCTGTTACA	1320
	ATGTTTAAAA AAAAAAAAAA AAAAAAAAAA AAAAAAYTCG AGGCTGGCTCC CGTACC	1376

30 (2) INFORMATION FOR SEQ ID NO: 15:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 502 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

5 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:

TAAAACAGTG CCTGCCTCAA AGGGAGGACT CAGTCAATAT CTGTTGAATG AATGAATGAA 60
 TAATTCCTTG GGTCAACGAA TGAATGGCTG AATGAATGAT TTCTCCTTTC CCTCGGCACT 120
 GTCTGGAGTC CCCAGGACAG GCATGGGCAG CAGTCGCTGG TCTGTGGCCT GTCCCACTGG 180
 10 ACTTGGGGTT CTCATGCTTG GTCTGGGCGG AGATCACCCA CCAGGCTCCC AGGTCGATCC 240
 TCTGCTCATG GGAARCTGCG TCCGGCCCNA GCTGCCAGAA CTCACTGCAS GGTGGAGGGA 300
 ARARCAGGRA CGATCTGCGA GCGCCTGAAC AGCGCACAA AGCCGAGGAG CCGCTGCTTA 360
 AAATGCAGGC GTTGAGAGGA GTTTCGCCTC CTTTTTTGAG TTGAATATGA GATTTCGAG 420
 CAGCCATGAC GAGTTGGGTT GGTGGAAGTG GGGAGTCCGT TCCTCAGTCA GATGGAGGAG 480
 15 GGGGTCCCCT TGGATCTCCT CT 502

(2) INFORMATION FOR SEQ ID NO: 16:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 425 base pairs

20 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

ATCTCTAGTG GTGGCTGCCG TCGCTCCAGA CAATCGGAAT CCTGCCTTCA CCACCATGGG 60
 25 CTGGCTTTTT CTAAAGGTTT TGTGGCGGG AGTGAGTTTC TCAGGATTTT TTTATCCTCT 120
 TGTGGATTTT TGCATCAGTG GGAAAACAAG AGGACAGAAG CCAAACTTTG TGATTATTTT 180
 GGCCGATGAC ATGGGGTGGG GTGACTGGGG AGCAAACTGG GCAGAAACAA AGGACACTGC 240
 CAACCTTGAT AAGATGGCTT CGGAGGGAAT GARGTGARTT TTGARATGCC ARGCCAGCTT 300
 TCTTTGGAAG TCTTACTCCC GTTCTTGAAA AGGGAAAGGG GCTTCAAAAG CACTTAARGA 360
 30 WTCATKGATG GACCCATGTG ATTTARTTAA TTTATTAATT AATTTGGTTT GGAARCCAGC 420

ATAGC

425

(2) INFORMATION FOR SEQ ID NO: 17:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 1316 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:

10	GGCACGAGGA GCTGGGGGAG CCTGAGGTGC GCTACGTGGC TGGCATGCAT GGGAACGAGG	60
	CCCTGGGGCG GGAGTTGCTT CTGCTCCTGA TGCAGTTCCT GTGCCATGAG TTCCTGCGAG	120
	GGAACCCACG GGTGACCCGG CTGCTCTCTG AGATGCGCAT TCACCTGCTG CCCTCCATGA	180
	ACCCTGATGG CTATGAGATC GCCTACCACC GGGGTTCAGA GCTGGTGGGC TGGGCCGAGG	240
	GCCGCTGGAA CAACCAGAGC ATCGATCTTA ACCATAATTT TGCTGACCTC AACACACCAC	300
15	TGTGGGAAGC ACAGGACGAT GGGAAGGTGC CCCACATCGT CCCCACCAT CACCTGCCAT	360
	TGCCCCCTTA CTACACCCTG CCCAATGCCA CCGTGGCTCC TGAAACGCGG GCAGTAATCA	420
	AGTGGATGAA GCGGATCCCC TTTGTGCTAA GTGCCAACCT CCACGGGGGT GAGCTCGTGG	480
	TGTCCTACCC ATTTCGACATG ACTCGCACCC CGTGGGCTGC CCGCGAGCTC ACGCCCACAC	540
	CAGATGATGC TGTGTTTCGC TGGCTCAGCA CTGTCTATGC TGGCAGTAAT CTGGCCATGC	600
20	AGGACACCAG CCGCCGACCC TGCCACAGCC AGGACTTCTC CGTGACACGGC AACATCATCA	660
	ACGGGGCTGA CTGGCACACG GTCCCCGGGA GCATGAATGA CTTGAGCTAC CTACACACCA	720
	ACTGCTTTGA GGTCACTGTG GAGCTGTCCT GTGACAAGTT CCCTCACGAG AATGAATTGC	780
	CCCAGGAGTG GGAGAACAAAC AAAGACGCCC TCCTCACCTA CCTGGAGCAG GTGCGCATGG	840
	GCATTGCAGG AGTGGTGAGG GACAAGGACA CGGAGCTTGG GATTGCTGAC GCTGTCATTG	900
25	CCGTGGATGG GATTAACCAT GACGTGACCA CGGCGTGGGG CGGGGATTAT TGGCGTCTGC	960
	TGACCCACAG GGAATACATG GTGACTGCCA GTGCCGAGGG CTACCATTTCA GTGACACGGA	1020
	ACTGTCGGGT CACCTTTGAA GAGGGCCCCCT TCCCCTGCAA TTTCGTGCTC ACCAAGACTC	1080
	CCAAACAGAG GCTGCGCGAG CTGCTGGCAG CTGGGGCCAA GTGCCCCCG GACCTTCGCA	1140
	GGCGCTGGA GCGGCTAAGG GGACAGAAGG ATTGATACCT GCGGTTTAAG AGCCCTAGGG	1200
30	CAGGCTGGAC CTGTCAAGAC GGGAAGGGGA AGAGTAGAGA GGGACGGACA AAGTGAGGAA	1260

AAGGTGCTCA TTAAAGCTAC CGGGCACCTT AAAAAAAAAA AAAAAAAAAA AAAAAA 1316

(2) INFORMATION FOR SEQ ID NO: 18:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 436 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:

10 AAAAAAATTC AATGGATATT ATGAAAATAA GAGAGTATTT CCAGAAAGTAT GGATATAGTC 60
CACGTGTCAA GAAAAATTCA GTACACGAGC AAGAAGCCAT TAACTCTGAC CCAGAGTTGT 120
CTAATTGTGA AAATTTTCAG AAGACTGATG TGAAAGATGA TCTGTCTGAT CCTCCTGTTG 180
CAAGCAGTTG TATTTCTGAG AAGTCTCCAC GTAGTCCACA ACTTTCAGAT TTTGGACTTG 240
AGCGGTACAT CGTATCCCAA GTTCTACCAA ACCCTCCACA GGCAGTGAAC AACTATAAGG 300
15 AAGAGCCCGT AATTGTAAAC CCACCTACCA AACAATCACT AGTAAAAGTA CTAAAACTC 360
CAAATGTGC ACTAAATGG ATGATTTTGA GTGTGTACTC CTAAATTAGA AACTTTGGT 420
ATCTCTGAAT ATACTA 436

(2) INFORMATION FOR SEQ ID NO: 19:

20 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 503 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:

TGTGCATATC CTGGGGAAAA AAATGGTACA TGTTTTAGAA ATTTTACTGT TTATAACAAT 60
GCAGGCAGTC AGTTTCCCGT TTCAAACACA GATAGATACA TGCAACACTC AAGATCCTGC 120
AGAGAGGCAG CCAGCATCTA TTGTTTAAAA AGGTTTCAAA AAGAATTCGG ATTGCTCKTT 180
TCTCTTTTGA ATCTGTGTGC CAAATGACAG GGACCAATAT TCGTCTTCTT TTTCKGTAAA 240
30 AYTCAAGAAAG AMACATGAAA GAACCCAGAA TGCATTTCTT AAAGGATTT AGTGCAGTTA 300

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TTTTAAATAA TTTATGCACG CACACACACA TACATATATC CCCCAGACTAC ATATTTTTC 360
 CCTTTTFACT TGTGTGCAAT CAGTAGCTAC AATGACTGAA ATCCACTTCT TTGGGACTGT 420
 GACATTTAAG CAAATCTTGT NTCTAGAAAN CGAAATGCCA NANTCTCGCA CAAAGCTGCT 480
 CCGTCTGGGG CAACAAATCC ACA 503

5

(2) INFORMATION FOR SEQ ID NO: 20:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 358 base pairs

(B) TYPE: nucleic acid

10

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

GGGCTGTCTC CCCAGTAGTA ACTTGCTGGC CCTGCCCTTG AAGTGGGGAA ACTGTGAAGG 60
 GCTCCTTGAT CAAGCTTGTC CTCMTTCTT ACCTCTTCCT CTCMTCTGTT TCCGCTGCAG 120
 15 CTGAACAGGC CAGCAGGCAA CCTGCCATGG GGTCTCTGCTC CAAGAACCGG TCCTTCTTCT 180
 GGATGACTGG GCTCCTGGTA TTCATCAGCC TCCTCCTCAG TGAGTGGCAG GGTCCCTGGG 240
 AAGGGAGGGC AATTGGAGAG GGCTGGGCTA GCTGGGCTCT GACCAACGGG TGGGCTGTTC 300
 AACTTCTGAT GTCTTTGGGC AACAAACACAG AAAAACACTC TGTATGATT TACGAAAN 358

20

(2) INFORMATION FOR SEQ ID NO: 21:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1926 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

25

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:

AGTGAAGGGA GCTGGCCGTG CGACTGGGCT TCGGGCCCTG TGCCAGAGGA GCANGCCTTC 60
 CTGAGCAGGA GGAAGCAGGT GGTGGCCCGG GCCTTGAGGC AGGCCCTTCCA GCTGGATGGA 120
 GACCTGCAGG AGGATGAGAT CCCAGTGGTA GCTATTATG CCACTGCTGG TGGGATCCGG 180
 30 GCAATGACTT CCCTGTATGG GCAGCTGGCT GGCTGAAGG AGCTGGGCT CTTGGATTGC 240

	KTCTCCTACA TCACCGGGGC CTCGGGCTCC ACCTGGGCCT TGGCCAACCT TTATAAGGAC	300
	CCAGAGTGGT CTCAGAAGGA CCTGGCAGGG CCCACTGAGT TGCTGAAGAC CCAGGTGACC	360
	AAGAACAAGC TGGGTGTGCT GGCCCCCAGC CAGCTGCAGC GGTACCGGCA GGAGCTGGCC	420
	GAGCGTGCCC GCTTGGGCTA CCCAAGCTGC TTCACCAACC TGTGGGCCCT CATCAACGAG	480
5	GCGCTGCTGC ATGATGAGCC CCATGATCAC AAGCTCTCAG ATCAACGGGA GGCCCTGAGT	540
	CATGGCCAGA ACCCTCTGCC CATCTACTGT GCCCTCAACA CCAAAGGGCA GAGCCTGACC	600
	ACTTTTGAAT TTGGGGAGTG GTGCGAGTTC TCTCCCTACG AGGTCGGCTT CCCCAAGTAC	660
	GGGGCCTTCA TCCCTCTGA GCTCTTTGGC TCCGAGTTCT TTATGGGGCA GCTGATGAAG	720
	AGGCTTCCTG AGTCCCGCAT CTGCTTCTTA GAAGGTATCT GGAGCAACCT GTATGCAGCC	780
10	AACCTCCAGG ACAGCTTATA CTGGGCCTCA GAGCCCAGCC AGTTCTGGGA CCGCTGGGTC	840
	AGGAACCAGG CCAACCTGGA CAAGGAGCAG GTCCCCCTTC TGAAGATAGA AGAACCACCC	900
	TCAACAGCCG GCAGAATAGC TGAGTTTTTC ACCGATCTTC TGACGTGGCG TCCACTGGCC	960
	CAGGCCACAC ATAATTTCTT GCGTGGCCTC CATTTCCACA AAGACTACTT TCAGCATCCT	1020
	CACTTCTCCA CATGGAAAGC TACCACTCTG GATGGGCTCC CCAACCAGCT GACACCCCTG	1080
15	GAGCCCCACC TGTGCCTGCT GGATGTTGGC TACCTCATCA ATACCAGCTG CCTGCCCCTC	1140
	CTGCAGCCCA CTCGGGACGT GGACCTCATC CTGTCA TTGG ACTACAACCT CCACGGAGCC	1200
	TTCCAGCAGT TGCAGCTCCT GGGCCGGTTC TGCCAGGAGC AGGGGATCCC GTTCCCACCC	1260
	ATCTCGCCCA GCCCCGAAGA GCAGCTCCAG CCTCGGGAGT GCCACACCTT CTCCGACCCC	1320
	ACCTGCCCCG GAGCCCCGTC GGTGCTGCAC TTTCTCTGG TCAGCGACTC CTTCCGGGAG	1380
20	TACTCGGCCC CTGGGTCCG GCGGACACCC GAGGAGGCGG CAGCTGGGGA GGTGAACCTG	1440
	TCTTCATCGG ACTCTCCCTA CCACTACAG AAGGTGACCT ACAGCCAGGA GGACGTGGAC	1500
	AAGCTGCTGC ACCTGACACA TTACAATGTC TGCAACAACC AGGAGCAGCT GCTGGAGGCT	1560
	CTGCGCCAGG CAGTGCAGCG GAGGCGGCAG CGCAGGCCCC ACTGATGGCC GGGGCCCTG	1620
	CCACCCCTAA CTCTCA TTCA TTCCCTGGCT GCTGAGTTGC AGGTGGGAAC TGTCAACACG	1680
25	CAGTGCTTNC AGAGCCTCGG GCTCAGGTGG CACTGTCCCA GGGTCCAGGC TGAGGGCTGG	1740
	GAGCTCCCTT GCGCCTCAGC AGTTTGAGT GGGTAAGGA GGGCAAGCCC ATTTGTGTAA	1800
30	TCACCCAAAA CCCCCGGCC TGTGCCTGTT TTCCCTCTTG CGCTACCTTG AGTAGTTGGA	1860
	GCACTTGATA CATCACAGAC TCATACAAAT GTGAGGCGCT GAGAAAAAAA AAAAAAAAAA	1920
	ACTCGA	1926

(2) INFORMATION FOR SEQ ID NO: 22:

5

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1224 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

10

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

15 CCGCCGAAGC TCCGTCCCGC CCGCGGCCGG CTCCGCCTCA CCTCCCGGCC GCGGCTGCCC 60
TCTGCCCGGG TTGTCCAAGA TGGAGGGCGC TCCACCGGGG TCGCTCGCCC TCCGGCTCCT 120
GCTGTTCGTG GCGCTACCCG CCTCCGGCTG GCTGACGACG GGCGCCCCCG AGCCGCCGCC 180
20 GCTGTCCGGA GCCCCACAGG ACGGCATCAG AATTAATGTA ACTACACTGA AAGATGATGG 240
GGACATATCT AAACAGCAGG TTGTTCTTAA CATAACCTAT GAGAGTGGAC AGGTGTATGT 300
AAATGACTTA CCTGTAAATA GTGGTGTAA CCGAATAAGC TGTCAGACTT TGATAGTGAA 360
25 GAATGAAAAT CTTGAAAATT TGGAGGAAAA AGAATATTTT GGAATTGTCA GTGTAAGGAT 420
TTTAGTTCAT GAGTGGCCTA TGACATCTGG TTCCAGTTTG CAACTAATG TCATTCAAGA 480
30 AGAGGTAGTA GAGATTGATG GAAAACAAGT TCAGCAAAG GATGTCACTG AAATTGATAT 540
TTTAGTTAAG AACCGGGGAG TACTCAGACA TTCAAACCTAT ACCCTCCCTT TGAAGAAAAG 600
CATGCTCTAC TCTATTTCTC GAGACAGTGA CATTTTATTT ACCCTTCCTA ACCTCTCCAA 660
35 AAAAGAAAGT GTTAGTTCAC TGCAAACCAC TAGCCAGTAT CTTATCAGGA ATGTGGAAAC 720
CACTGTAGAT GAAGATGTTT TACCTGGGCA AGTTACCTGA AACTCCTCTC AGAGCAGAGC 780
40 CGCCATCTTC ATATAAGGTA ATGTGTCAGT GGATGGAAAA GTTTAGAAAA GATCTGTGTA 840
GGTTCTGGAG CAACGTTTTT CCAGTATCTT TTCAGTTTTT GAACATCATG GTGGTTGGAA 900
TTACAGGAGC AGCTGTGGTA ATAACCATCT TAAAGGTGTT TTTCCCAGTT TCTGAATACA 960
45 AAGGAATTCT TCAGTTGGAT AAAGTGGACG TCATACCTGT GACAGCTATC AACTTATATC 1020
CAGATGGTCC AGAGAAAAGA GCTGAAAACC TTGAAGATAA AACATGTATT TAAAACGCCA 1080
50 TCTCATATCA TGGACTCCGA AGTAGCCTGT TGCCTCCAAA TTGCCACTT GAATATAATT 1140
TTCTTTAAAT CGTTAAGAAT CAGTTTATAC ACTAGAGAAA TTCTTAAAT CTAAGACTGC 1200
CTGAAAATTG ACCTTTACAG TGCC 1224
55

(2) INFORMATION FOR SEQ ID NO: 23:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 694 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

10 GGCACGAGTC TTATTGTGCA CTGTAGCCTG AATCCCCCAG GGTAATTAAT ATGAAGTGCA 60
AAAAGTTGAA TGTTCCAGTC TAAAAGGCAG TGGGAGAAAT TACATAGCAT GGAAATAATA 120
15 AAATGAAGTC TTATTAATGA GAACGAGGCT CTTGCAGTGG CAAGTTCTGC TGGTCACCCG 180
ATGGGGATGG GAGCCTTTCA AGCTTTTTTT TGGGTAATAC TCACAGTTTC CAACGTCTGT 240
20 GTACTTTTCA AAATGAGCTT GTTCTTCCTT CTGACACTCA TCTCAAAGCT CCATGGTGAC 300
GCAGAGGTCT GTTGAAGGTC ACAGGTCCTC GCTTGCATTG GCATACGGTC CTGTAGCATC 360
ACTTGTTAGC CCACTGCTGC TTGAAGGAAC TAAGAGTATT CAGGGATAGA GAGCTGAAAA 420
25 TAGGATTAAT TCCTTCCTTT TGA CTCTCTCCC CTCAAGATGT CCTTGCTTTG GTCTGAAAAC 480
CTCTCCTGAC AACTTTTGCC CAAAGCAAAC CATCTGCCTT TTCTGAACTC TGAGTGAATA 540
TATTAGCATC TTCCCTCTCG AGCCCTCGTA CTGCCANGTT TGTGTGTTTG TTTGTTTCCA 600
30 AGAGACTGTG TCTTGCTCTG TCACCCAGGA GTTTGAAACC AGCCTGGCAA CATAGCAAGA 660
CCCTATCTCT ACAAAAAAAAA AAAAAAAAAA AAAA 694
35

(2) INFORMATION FOR SEQ ID NO: 24:

(i) SEQUENCE CHARACTERISTICS:

- 40 (A) LENGTH: 796 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:

ATGAGCGGCG GTTGGATGGC GCAGGTTGGA GCGTGGCGAA CAGGGGCTCT GGGCCTGGCG 60
50 CTGCTGCTGC TGCTCGGCCT CGGACTAGGC CTGGAGGCGC GCGGAGCCCG CTTTCCACCC 120
CGACCTCTGC CCAGGCCGCA CCCGAGCTCA GGCTCGTGCG CACCCACCAA GTTCCAGTGC 180
55 CGCACCAGTG GCTTATGCGT GCCCCTCACC TGGCGCTGCG ACAGGACTTG TACTGCAGCG 240
ATGGCAGCGA TGAGGAGGAG TGCAGGATTG AGCCATGTAT CCAGAAAGCG CAATGCCCCAC 300
CGCCCCCTGG CCTCCCTGTC CCCTGCACCG GCGTCAGTGA CTGCTCTGCG GGAAGTGACA 360
60 AGAAACTGCG CAACTGCAGC CGCCTGGCCT GCCTAGCAGS GRAGSKCMCG WKGCACGCTG 420

175

AGCGATGACT GCATTCCACT CACGTGGCGC TCGACGGCC ACCCAGACTG TCCCGACTCC 480
 AGCGACGAGC TCGGCTGTGG AACCAATGAG ATCCTCCCGG AAGGGGATGC CACAACCATG 540
 5 GGGCCCCCTG TGACCCTGGA GAGTGTCAAC TCTCTCAGGA ATGCCACAAC CATGGGGCCC 600
 CCTGTGACCC TGGAGAGTGT CCCCTCTGTC GGAATGCCA CATCCTCCTC TGCCGGAGAC 660
 10 CAGTCTGGAA GCCCAACTGC CTATGGGGTT ATGTCAGCTG CTGCGGTGCT CAGTGCAAGC 720
 CTGGTCACCG CCACCCTCCT CCTTTTGTCC TGGCTCCGAG CCCAGGAGCG CCTCCGCCCA 780
 CTGGGGTTAC TGGTGG 796
 15

20 (2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 662 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 25 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:
 TAATTTCGGCA CGAGGCTGTG GTGGAGAAGG ACGTGCCGTG CCGCTGGGTT CTGAGCCGGA 60
 30 GTGGTCGGTG GGTGGGATGG AGGCGACCTT GGAGCAGCAC TTGGAAGACA CAATGAAGAA 120
 TCCCTCCATT GTTGGAGTCC TGTGCACAGA TTCACAAGGA CTTAATCTGG GTTGCCGCGG 180
 35 GACCCTGTCA GATGAGCATG CTGGAGTGAT ATCTGTTCTA GCCCAGCAAG CAGCTAAGCT 240
 AACCTCTGAC CCCACTGATA TTCTGTGGT GTGTCTAGAA TCAGATAATG GGAACATTAT 300
 GATCCAGAAA CACGATGGCA TCACGGTGGC AGTGCACAAA ATGGCCTCTT GATGCTCATA 360
 40 TCTGTTCTTC AGCAGCCTGT CATAGGAACCT GGATCCTACC TATGTTAATT ACCTTATAGA 420
 ACTACTAAAG TTCCAGTAGT TAGGCCATTC ATTTAATGTG CATTAGGCAC TTTTCTGTTT 480
 45 ATTTAAGAGT CAATTGCTTT CTAATGCTCT ATGGACCGAC TATCAAGATA TTAGTAAGAA 540
 AGGATCATGT TTTGAAGCAG CAGGTCCAGG TCACTTTGTA TATAGAATTT TGCTGTATTC 600
 AATAAATCTG TTTGGAGGAA AAAAAAAAAA AAAAAAATTA CTTGGGNCCTG ACAAGGGAAT 660
 50 TC 662

55 (2) INFORMATION FOR SEQ ID NO: 26:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1105 base pairs
 60 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:

```

5      CCTGATCCTC TCTTTTCTGC AGTTCAAGGG AAAGACGAGA TCTTGACAAA GGCACCTCTGC      60
      TTCTGCCCTT GGCTGGGGAA GGGTGGCATG GAGCCTCTCC GGCTGCTCAT CTTACTCTTT      120
10     GTCACAGAGC TGTCCGGAGC CCACAACACC ACAGTGTTCG AGGGCGTGGC GGGCCAGTCC      180
      CTGCAGGTGT CTTGCCCTTA TGA CTCCATG AAGCACTGGG GGAGGCGCAA GGCCTGGTGC      240
      CGCCAGCTGG GAGAGAAGGG CCCATGCCAG CGTGTGGTCA GCACGCACAA CTTGTGGCTG      300
15     CTGTCCCTTC TGAGGAGGTG GAATGGGAGC ACAGCCATCA CAGACGATAC CCTGGGTGGC      360
      ACTCTACCA TTACGCTGCG GAATCTACAA CCCCATGATG CGGGTCTCTA CCAGTGCCAG      420
20     AGCCTCCATG GCAGTGAGGC TGACACCTC AGGAAGGTCC TGGTGGAGGT CCTCGCAGAC      480
      CCCCTGGATC ACCGGGATGC TGGAGATCTC TGGTTCCCCG GGGAGTCTGA GAGCTTCGAG      540
      GATGCCCATG TGGAGCACAG CATCTCCAGG AGCTCTTCKT AGGAAAGGCC GCAAATTTCC      600
25     ATTCCTTCCC CTCTTGCCCTA TCYTTCTCCT CCAAGAYCTG CATCTTTCTC ATCAAGATTC      660
      TAGCAGCCAG CGCCCTCTGG GCTGCAGCCT GGCATGGACA GAAGCCAGGG ACACATCCAC      720
30     CCAGTGAAGT GGA CTGTGGC CATGACCCAG GGTATCAGCT CCAAACCTCTG CCAGGGCTGA      780
      GAGACACGTG AAGGAAGATG ATGGGAGGAA AAGCCCAGGA GAAGTCCCAC CAGGACCAG      840
      CCCAGCCTGC ATACTTGCCA CTTGGCCACC AGGACTCCTT GTTCTGCTCT GGCAAGAGAC      900
35     TACTCTGCCT GAACACTGCT TCTCCTGGAC CCTGGAAGCA GGGACTGGTT GAGGGAGTGG      960
      GGAGGTGGTA AGAACACCTG ACAACTTCTG AATATTGGAC ATTTTAAACA CTTACAAATA      1020
40     AATCCAAGAC TGTCATATTT AAAAAAAAAA AAAAAAAMA AAARRRRRRR CCCGGTACCC      1080
      AATTCGCCCT ATAGTGAGTC GTATA      1105

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45

(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:

```

50     (A) LENGTH: 1017 base pairs
      (B) TYPE: nucleic acid
      (C) STRANDEDNESS: double
      (D) TOPOLOGY: linear

```

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

```

      CTCGCCCTGGG CTGTTTCCCC GCTTCATTC TCCCGACTTA TTTTCTTCTC CTGGGCTTTC      60
      CGAGGTGCTT TCGCCGCTGT CCCCACCACT GCAGCCATGA TTTTCTTAC GGACACGCAG      120

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60

177

	AAAATTGGAA TGGGATTAAC AGGATTTGGA GTGTTTTTCC TGTTCMTTGG AATGATTCTC	180
	TTTTTTGACA AAGCACTACT GGCTATTGGA AATGTTTTAT TTGTAGCCGG CTGGGCTTTT	240
5	GTAATTGGTT TAGAAAGAAC ATTCAGATTC TTCTTCCAAA AACATAAAAT GAAAGCTACA	300
	GGTTTTTTTC TGGGTGGTGT ATTTGTAGTC CTATTGGTT GGCCTTTGAT AGGCATGATC	360
10	TTCGAAATTT ATGGATTTTT TCTCTGTTC AGGGGCTTCT TTCCTGTCGT TGTGGCTTT	420
	ATTAGAAGAG TGCCAGTCCT TGGATCCCTC CTAAATTTAC CTGGAATTAG ATCATTTGTA	480
	GATAAAGTTG GAGAAAGCAA CAATATGGTA TAACAACAAG TGAATTTGAA GACTCATTTA	540
15	AAATATTGTG TTATTTATAA AGTCATTTGA AGAATAITCA GCACAAAATT AAATTACATG	600
	AAATAGCTTG TAATGTTCTT TACAGGAGTT TAAAACGTAT AGCCTACAAA GTACCAGCAG	660
20	CAAATTAGCA AAGAAGCAGT GAAAACAGGC TTCTACTCAA GTGAACTAAG AAGAAGTCAG	720
	CAAGCAAAC T GAGAGAGTG AAATCCATGT TAATGATGCT TAAGAACTC TTGAAGGCTA	780
	TTTGTGTTGT TTTTCCACAA TGTGCGAAAC TCAGCCATCC TTAGAGAACT GTGGTGCCTG	840
25	TTTCTTTTCT TTTTATTTTG AAGGCTCAGG AGCATCCATA GGCATTTGCT TTTTAGAAAT	900
	GTCCACTGCA ATGGCAAAAA TATTTCCAGT TGCACGTAT CTCTGGAAGT GATGCATGAA	960
30	TTCGATTGGA TTGTGTCATT TTAAAGTATT AAAACCAAGG GAAACCCCAA AAAAAA	1017

35 (2) INFORMATION FOR SEQ ID NO: 28:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 391 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

45	CCCTGGAAAG AGGAACTGAT GTTTGAGGGG ACAGATGTGG GTCACTTTCC CTGGCAGTGC	60
	CCTCTAGCCT TGCTGCCTTG GCTTCTGAC CCCTTCCAGG CTTCAGGGGC CTGGGAGATC	120
	TCATGCCTCA GCCCAGGAAA CATTTAATAG GGAAAGCAGA GACATGTCAT GTCAGCCCCA	180
50	CAGACAAGAA TTTCTAGAGC ACTTGTCTTG TTGTTCTTGG CCCCAGACAT ACTCAGTCTG	240
	GGCCATGGAA TCCATCCAAT AAACACAGCA ACACCTATG MTACTGACCA AGCAAAGCTT	300
55	CCCCCTGGTA CCAAAGAGCT AAATCATGAC CAAAGTGTGA CATGAATGTA ACTGAAATGC	360
	GGTTAGTTG CTCAATGTAT GCAAAGTCCC A	391

60

(2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 1139 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

10 GGTGATATCT TCATAGTGGG CTATTACAGG CAGGAAAATG TTTTAACTGG TTTACAAAAT 60
CCATCAATAC TTGTGTCATT CCTGTAAAA GGCAGGAGAC ATGTGATTAT GATCAGGAAA 120
15 CTGCACAAAA TTATTGTTTT CAGCCCCCGT GTTATTGTCC TTTTGAACGT TTTTTTTTTT 180
ATTAAAGCCA AATTTGTGTT GTATATATTC GTATTCCATG TGTTAGATGG AAGCATTTCC 240
TATCCAGTGT GAATAAAAAG AACAGTTGTA GTAAATTATT ATAAAGCCGA TGATATTTCA 300
20 TGGCAGGTTA TTCTACCAAG CTGTGCTTGT TGGTMTTTC CATGACTGTA TTGCTTTTAT 360
AAATGTACAA ATAGTTACTG AAATGACGAG ACCCTTGTTT GCACAGCATT AATAAGAACC 420
25 TTGATAAGAA CCATATCTCG TTGACAGCCA GCTCACAGTT TCTTGCCTGA AGCTTGGTGC 480
ACCCTCCAGT GAGACACAAG ATCTCTCTTT TACCAAAGTT GAGAACAGAG CTGGTGGATT 540
AATTAATAGT CTTGATATC TGGCCATGGG TAACCTCATT GTAACATCA TCAGAATGGG 600
30 CAGAGATGAT CTTGAAGTGT CACATACACT AAAGTCCAAA CACTATGTCA GATGGGGGTA 660
AAATCCATTA AAGAACAGGA AAAAATAATT ATAAGATGAT AAGCAAATGT TTCAGCCCCA 720
35 TGTCAACCCA GTTAAAAAAA AATTAATGC TGTGTAAAAT GGTGAATTA GTTTGCAAAC 780
TATATAAAGA CATATGCAGT AAAAAGTCTG TTAATGCACA TCCTGTGGGA ATGGAGTGTT 840
CTAACCAATT GCCTTTTCTT GTTATCTGAG CTCTCCTATA TTATCATACT CAGATAACCA 900
40 AATTAAAGA ATTAGAATAT GATTTTAAAT ACACCTAACA TTAAACTCTT CTAACTTTCT 960
TCTTCTGTG ATAATTCAGA AGATAGTTAT GGATCTTCAA TGCTCTGAG TCATTGTTAT 1020
45 AAAAAATCAG TTATCACTAT ACCATGCTAT AGGAGACTGG GCAAAACCTG TACAATGACA 1080
ACCCTGGAAG TTGCTTTTTT TAAAAAATA ATAAATTCTT TAAATCAAAA AAAAAAAAAA 1139

50

(2) INFORMATION FOR SEQ ID NO: 30:

(i) SEQUENCE CHARACTERISTICS:

- 55 (A) LENGTH: 465 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

60

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

5 CCACGCGTCC GCGGACGCGT GGGGAAGGTT TGTGCCAGTA GACATTATGT TACTAAATCA 60
 GCACTTTTAAA ATCTTTGGTT CTCTAATTCA TATGAATTTG CTGTTTGCTC TAATTTCTTT 120
 GGGCTCTTCT AATTTGAGTG GAGTACAATT TTGTTGTGAA ACAGTCCAGT GAAACTGTGC 180
 AGGGAAATGA AGGTAGAATT TTGGGAGGTA ATAATGATGT GAAACATAAA GATTTAATAA 240
 10 TTACTGTCCA ACACAGTGGA GCAGCTTGTC CACAAATATA GTAATTACTA TTTATTGCTC 300
 TAAGGAAGAT TAAAAAAGA TAGGGAAAAG GGGGAAACTT CTTTGAAAAA TGAAACATCT 360
 GTTACATTAA TGTCTAATTA TAAATTTTA ATCCTTACTG CATTTCTTCT GTTCCTACAA 420
 15 ATGTATTAAA CATTCAGTTT AACTGGTAAA AAAAAAAAAA AAAAA 465

20

(2) INFORMATION FOR SEQ ID NO: 31:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 702 base pairs
 25 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:

30 GCAACAAGCG GCCACCTTC CTGAAGATCA AGAAGCCACT GTCGTACCGC AAGCCCATGG 60
 ACACGGACCT GGTGTACATC GAGAAGTCGC CCAACTACTG CGAGGAGGAC CCGGTGACCG 120
 35 GCAGTGTGGG CACCCAGGGC CGCGCCTGCA ACAAGACGGC TCCCCAGGCC AGCGGCTGTG 180
 ACCTCATGTG CTGTGGGCGT GGCTACAACA CCCACCAGTA CGCCCGCGTG TGGCAGTGCA 240
 ACTGTAAGTT CCACTGGTGC TGCTATGTCA AGTGCAACAC GTGCAGCGAG CGCACGGANG 300
 40 ATGTACACGT GCAAGTGAGC CCCGTGTGCA CACCACCCTC CCGCTGCAAG TCAGATTGCT 360
 GGGAGGACTG GACCGTTTCC AAGCTGCGGG CTCCCTGGCA GGATGCTGAG CTTGTCTTTT 420
 45 CTGCTGAGGA GGGTACTTTT CCTGGGTTTC CTGCAGGCAT CCGTGGGGA AAAAAATCT 480
 CTCAGAGNCC TCAACTATTC TGTCCACAC CCAATGCTGS TCCACCCTCC CCCAGACACA 540
 GCCCAGGTCC CTCCGCGGCT GGAGCGAAGC CTTCTGCAGC AGGAACTCTG GACCCCTGGG 600
 50 CCTCATCACA GCAATATTTA ACAATTTATT CCTGATAAAA ATAATATTAA TTTATTTAAT 660
 TAAAAAGAAT TCTTCCAAA AAAAAAAAAA AAAAAACNT CG 702

55

(2) INFORMATION FOR SEQ ID NO: 32:

60 (i) SEQUENCE CHARACTERISTICS:

180

- (A) LENGTH: 1142 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 32:

```
CGGCACGAGG AAGAAATGGC AGAGACTGGA ATCTCTCTTC ATGAAAAAAT GCAGCCCCCTT      60
10 AACTTCAGTT CGACAGAGTG CAGCTCCTTC TCTCCACCCA CCACAGTGAT TCTCCTTATC      120
CTGCTGTGCT TTGAGGGCCT GCTCTTCCTC ATTTTCACAT CAGTGATGTT TGGGACCCAG      180
15 GTGCACTCCA TCTGCACAGA TGAGACGGGA ATAGAACAAT TGAAAAAGGA AGAGAGAAGA      240
TGGGCTAAAA AAACAAAATG GATGAACATG AAAGCCGTTT TTGGCCACCC CTTCTCTCTA      300
GGCTGGGCCA GCCCCTTTGC CACGCCAGAC CAAGGGAAGG CAGACCCGTA CCAGTATGTG      360
20 GTCTGAAGGA CCCCACCCG CATGGCCACT CAGACACAAG TCCACACCAC AGCACTACCG      420
TCCCATCCGT TCTCATGAAT GTTTAAATCG AAAAAGCAAA ACAACTACTC TTAATACTTT      480
25 TTTTATGTCT CAAGTAAAT GGCTGAGCAT TGCAGAGARA AAAAAAGTC CCCACATTTT      540
ATTTTTTAAA AACCATCCTT TCGATTCTT TTGGTGACCG AAGCTGCTCT CTTTCTCTTT      600
TAAAATCACT TCTCTGGCCT CTGGTTTCTC TCTGCTGTCT GTCTGGCATG ACTAATGTAG      660
30 AGGGCGCTGT CTCGCGCTGT GCCCATCTTA CTAAGTGAAT GAGACATGAC GCTGTGCTGG      720
GATGGAATAG TCTGGACACC TGGTGGGGGA TGCATGGGAA AGCCAGGAGG GCCCTGACCT      780
35 TCCCCTGCC CAGGAGGCAG TGGCGGGCTC CCCGATGGGA CATAAACCT CACCGAAGAT      840
GGATGCTTAC CCCTTGAGGC CTGAGAAGGG CAGGATCAGA AGGGACCTTG GCACAGCGAC      900
CTCATCCCCC AAGTGGACAC GGTTCGCTG CTAAGTGCAT AAGCAATTGC CTGCCTTGTA      960
40 CTTTATGGGC TTGGGGTGTG TAGAATGATT TTGCGGGGGA GTGGGGGAGA AAGATGAAAG      1020
AGGTCTTATT TGTATCTGA ATCAGCAATT ATATCCCTG TGATTATTTG GAAGAGTGTG      1080
TAGGAAAGAC GTTTTCCAG TTCAAAATGC CTTATACAAT CAAGAGGAAA AAAAAAAAAA      1140
45 AG                                                                                   1142
```

50

(2) INFORMATION FOR SEQ ID NO: 33:

(i) SEQUENCE CHARACTERISTICS:

55

- (A) LENGTH: 928 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

60

GGCACGAGGT CTAATGAGGG CTCTCTTGTT TGCTAGAGAT GAGAGAAATG TATACTAATC 60
 ATTTTAATTT GTACTTAAAA TACATTTTAC TAATCATATT GATTTTAAAT ATGACAAATT 120
 5 CTTCTAGTAG ATACTAATCT TTCTTGTTTA TCATATGTG CTAGAGAAGC CTAGGTAAAA 180
 ATGGGTTCCTA CCTAGTCTGT TTGTATAACA CCTTCCCCCG TCCCCTCTCC ATCCCTGCCA 240
 ATTGGGCTCT ATGCATATG ACAAGCAAAT AAGAAAACCT TAGGTTCTTG TATTTGAATT 300
 10 TCCAAAACAA TAAAAGGTTT TGAACAAGA TTTGCATTCA AGAAGAGGCA GAAATTTTGT 360
 CTTATCTTTT TATCATTTTG TGAACCTGTG TTTCTCTGTA TGCTTAGAAA ATTTACACAC 420
 15 AAGGAATGTT TGAAAAAGTG AGAATTTTAG AGTGCTTGGG TGGTTTTTAT TTGGTCAGTG 480
 CTGATGTGTT AGGTGTTTAG GGAAATAATG CTTCAGGACC TTTTGTACAA CACAGCTTCA 540
 TGAATGACTG GGGGATATTT ATGTTTGTGC TGAGAAAAGG GAGGGAGTGG GCAGGTGGA 600
 20 GTGGGGACCT TTCCATTGAA AGCAGTGCAG TCAGCTGTTT CGTAGATGCA TTTTCTCTTT 660
 ATGCTTGTA CATTGTTCTT GTGTCCATAA TTGACTGAAA TGTCAGCTC CAGGAATGCA 720
 25 AGGCATTAT CAGGTGACCA GAAGTAGAAC CTTGTTGATT ATGAAATGGA AGAATAATGT 780
 CAAGGTAGTG GGGGTAAAAT GACAAATAAG ATTTTACTGG TGAATTTCCA TGCTTAGTAT 840
 GTACATTAAC CTCTTTTAA GTTGCATGTT AATCTGGTAT AACGTATTGT GTCTGGTTTA 900
 30 TGCTTTGAGT AAAAAAAAAA AAAAAAAA 928

35

(2) INFORMATION FOR SEQ ID NO: 34:

(i) SEQUENCE CHARACTERISTICS:

40

- (A) LENGTH: 773 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:

45

GGCACGAGTT CTGGCCTCTC ATTCCTTAC ACTCTGACAT GAATGAATTA TTATTATTTT 60
 TCTTTTCTT TTTTTTTTTT ACATTTTGTA TAGAAACAAA TTCATTTAAA CAAACTTATT 120
 50 ATTATTATTT TTTACAAAAT ATATATATGG AGATGCTCCC TCCCCCTGTG AACCCCCAG 180
 TGCCCCCTG GGGCTGAGTC TGTGGGCCCA TTCGGCCAAG CTGGATTCTG TGTACCTAGT 240
 ACACAGGCAT GACTGGGATC CCGTGTACCG AGTACACGAC CCAGGTATGT ACCAAGTAGG 300
 55 CACCCTTGGG CGCACCCACT GGGGCCAGGG GTCGGGGGAT GTTGGGAGCC TCCTCCCCAC 360
 CCCACCTCCC TCACTTCACT GCATTCCAGA TTGGACATGT TGCATAGCCT TGCTGGGGAA 420
 60 GGGCCCACTG CCAACTCCCT CTGCCCCAGC CCCACCTTG GGCATCTCCC TTTGGGAACT 480

5 AGGGGGCTGC TGGTGGGAAA TGGGAGCCAG GGCAGATGTA TGCATTCCCTT TATGTCCCTG 540
 TAAATGTGGG ACTACAAGAA GAGGAGCTGC CTGAGTGGTA CTTTCTCTTC CTGGTAATCC 600
 TCTGGCCCAG CCTTATGGCA GAATAGAGGT ATTTTTAGGC TATTTTGTGA ATATGGCTTC 660
 TGGTCAAAAT CCCTGTGTAG CTGAATTCCT AAGCCCTGCA TTGTACAGCC CCCCCTCCC 720
 10 CTCACCACCT AATAAAGGAA TAGTTAACAC TCAAAAAAAA AAAAAAAAAA AAA 773

15 (2) INFORMATION FOR SEQ ID NO: 35:

(i) SEQUENCE CHARACTERISTICS:

20 (A) LENGTH: 453 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:

25 TAAAATGTTA CACGCTTGTC ATATTCCAGG CACTGCACTA TGTATGCCGT TTATCAACAG 60
 TTAGCTCAGC TAACCCTCAT GGTAACCTTG TTAGCCCCGA TTTTGCCAGA TGAGCAAAGT 120
 GAGGTTTTTG AGGCCTTAAG TAACTTGCCC AAGGTCACGT GGCTGGGAAG TAACTCTCCC 180
 30 AGTTCTGAGA TGCCCCGAGCC TGGACGCTTT GTCATTGTAC ACCATCAACT CAGTGCTGCC 240
 AGTCATTCCA GCAGCCAGCT AGCGTAGTCA AGGTTTCTCC ACCTTAGCAC TGTTGACATT 300
 35 TCGAGCCAGA TAATTCTCTG TGGTGAGGAG CTGTCCTATG CCTTGTAGGA TATACAACAG 360
 CATCYTGGCT TTACCCACCA GATGYTGGAA CACCTCCCCA GTCGTGACAG CCCAAAATGT 420
 CTATAGACGT TGCCACGTAT ACCCAGGGGT TCC 453
 40

45 (2) INFORMATION FOR SEQ ID NO: 36:

(i) SEQUENCE CHARACTERISTICS:

50 (A) LENGTH: 459 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:

55 GTGACTGCCG CCGTGCCCGC AGCCATGTGG CCCCCGCTGT TGCTGCTGCT CCTGCTGCTC 60
 CCGGCCGCCG CGGTCCCCAC CGCCAAAGCC GCTCCCCAGC CTGATGCTAA CACCCAGGAA 120
 GGCCTTCAGA ACCTGCTCCA AGGAGTCGGG GCTGGCGGAG ACGGAGAGCT CCGGGCAGAC 180
 60 TCACACCTGG CCCCAGGGCTC TGGCTGTATT GATGGGGCTG TGGTGGCCAC GCGACCAGAA 240

183

AGCCGGGGAG GAAGACCTGC GGTTCCTGA GAGGCGTCCA GGGCTGCAGG CCACGGCGAC 300
 AGGCTCCGGG GAACATGGGG CTTTCCCTGT CCACTCCCAA GGAGTGTGGG CCTCAACGCA 360
 5 TTGGCAGGGG ACGGCCGTGT GCCCTCTYCA GACCCACCC CCAGATGCAT TTATTAGAAA 420
 TAATAAATTC TTTCTTAGCT AAAAAAAAAA AAAAAAAT 459

10

(2) INFORMATION FOR SEQ ID NO: 37:

15

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 509 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 37:

ATGAAATTTA CCACTCTCCT CTTCTTGGCA GCTGTAGCAG GGGCCCTGGT CTATGCTGAA 60
 25 GATGCCTCCT CTGACTCGAC GGTGCTGAT CCTGCCCAGG AAGCTGGGAC CTCTAAGCCT 120
 AATGAAGAGA TCTCAGGTCC AGCAGAACCA GCTTCACCCC CAGAGACAAC CACAACAGCC 180
 CAGGAGACTT CGGCGGCAGC AGTTCAGGGG ACAGCCAAGG TCACCTCAAG CAGGCAGGAA 240
 30 CTAAACCCCC TGAATCCAT AGTGGAGAAA AGTATCTTAC TAACAGAACA AGCCCTTGCA 300
 AAAGCAGGAA AAGGAATGCA CGGAGGCGTG CCAGGTGGAA AACAATTCAT CGAAAATGGA 360
 35 AGTGAATTG CACAAAAATT ACTGAAGAAA TTCAGTCTAT TAAACCATG GGCATGAGAA 420
 GCTGAAAAGA ATGGGATCAT TGGACTTAAA GCCTTAAATA CCCTGTAGC CCAGAGCTAT 480
 TAAACGAAA GCATCAAAA AAAAAAAAAA 509
 40

45

(2) INFORMATION FOR SEQ ID NO: 38:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 598 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

50

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38:

ATGTTGGGCT GTGGGATCCC AGCGCTGGGC CTGCTCCTGC TGGTGCAGCG CTCGGCAGAC 60
 55 GGAAATGGAA TCCAGGGATT CTTCTACCCA TGGAGCTGTG AAGGTGACAT ATGGGACCGG 120
 GAGAGCTGTG GGGGCCAGGC GGCCATCGAT AGCCCCAACC TCTGCCTGCG TCTCCGGTGC 180
 60 TGCTACCGCA ATGGGGTCTG CTACCACCAG CGTCCAGACG AAAACGTGCG GAGGAAGCAC 240

184

5 ATGTGGGCGC TGGTCTGGAC GTGCAGCGGC CTCCTCCTCC TGAGCTGCAG CATCTGCTTG 300
 TTCTGGTGGG CCAAGCGCCG GGACGTGCTG CATATGCCCG GTTTCCTGGC GGGTCCGTGT 360
 GACATGTCCA AGTCCGTCTC GCTGCTCTCC AAGCACCGAG GGACCAAGAA GACGCCGTCC 420
 ACGGGCAGCG TGCCAGTCGC CCTGTCCAAA GAGTCCAGGG ATGTGGAGGG AGGCACCGAG 480
 10 GGGGAAGGGA CGGAGGAGGG TGAGGAGACA GAGGCGAGG AAGAGGAGGA TTAGGGGAGT 540
 CCCCCGGGGA CTGGTCAATA CAGATACGGT GGACGGAAAA AAAAAAAAAA AAAAAAAAAA 598

15

(2) INFORMATION FOR SEQ ID NO: 39:

20 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 454 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:

ATGGAGGCTG TTTTACAGT TTTTTTTTTT GTTGTGTTT TGTTTTAA GAATACAGAA 60
 GGAGCCAAGC TTTTTTGCAC TTTGTATCCA GCTGCAAGCT CAGGGCAGAG TCAAGGGCCT 120
 30 GGGTTGGAAA AACCTGACTC ACAGGAATGC ATAATTGACC CTTGCAGCTA CCCAATAGCC 180
 CTTGGAGCTG GCACTGAACC AGGCTGCAAG ATTGACTGCT CTAAAAACA CAAGGCCCTC 240
 35 TAGGCCTGGC AGGGATGTCC CTGTGCCCAG CACTGGGGGC TCGAAGACTG GTTCTAGCA 300
 CTACCGGTCA CGGCCATGTC GTCCTAGAAG GGTCCAGAAG ATTATTTTAC GTTGAGTCCA 360
 TTTTAAATGT TCTGATCACC TGACAGGGCA CCCCAAACCC CCAACTCCCA ATAAAAGCCG 420
 40 TGACGTTCCG AAAAAAAAAA AAAAAAAAAA AAAA 454

45

(2) INFORMATION FOR SEQ ID NO: 40:

50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 425 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:

GCTAAAGGCC ATTCCCTCCG CAGGGCATTG GCGTCGGGT GGGAGGGGAA AACGCATCTT 60
 GTTAATTATT TTTAATCTTA TTTATTGTAC ATACCTGGGG CAGGGGCTTG GGGAGGTGGA 120
 60 GGGGGRAGAA GGGTCCCCTC TCTCTGCCCC TCCCCTCCT TTTCTACGGC GATTGTCTG 180

185

5 TGTCTGGCCC CCACCCACTG MCCATCCCCC ATTGTTGTCT GGATGTGGTT CTATTTTPTA 240
 TCGGTCTCCT TTCCCCTCCT CCCCCTTTC GCCCCCGMCC CACCCCTGC TCCCCTACC 300
 CTTTGTCTCT TGCTCTTCT TGGGYTCTG TACAACCTAA CTTGTATACA CTGTGTACAC 360
 ACAACCAGYC WAACGCAAAA CCCAACGGCA AACACTTTAA AAAAAAAAAA AAAAAACTGG 420
 10 GGGGT 425

15 (2) INFORMATION FOR SEQ ID NO: 41:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2471 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:

25 GGCACGAGTA TGGCTTCCCG TGGACTCAGC CTCTTCCCG ANTCCTGGCA CGAGGGGGCT 60
 TCGCGTCTGT GCTTCCTGTG GCTGACGTCA TCTGGAGGAG ATTTGCTTTC TTTTCTCCA 120
 AAAGGGGAGG AAATTGAAAC TGAGTGGCCC ACATGGGAA GAGGGGAAAG CCCAGGGGTA 180
 30 CAGGAGGCCT CTGGGTGAAG GCAGAGGCTA ACATGGGGTT CGGAGCGACC TTGGCCGTTG 240
 GCCTGACCAT CTTGTGTCTG TCTGTCTGCA CTATCATCAT CTGCTTCACC TGCTCCTGCT 300
 35 GCTGCCTTTA CAAGACGTGC CGCCGACCAC GTCCGGTTGT CACCACCACC ACATCCACCA 360
 CTGTGGTGCA TGCCCCTTAT CCTCAGCCTC CAAGTGTGCC GCCCAGCTAC CCTGGACCAA 420
 GCTACCAGGG CTACCACACC ATGCCGCTC AGCCAGGGAT GCCAGCAGCA CCCTACCCAA 480
 40 TGCAGTACCC ACCACCTTAC CCAGCCCAGC CCATGGGCC ACCGGCCTAC CACGAGACCC 540
 TGGCTGGAGA GCAGCCGCGC CCTACCCCGC CAGCCAGCCT CCTTACAACC CGGCCTACAT 600
 45 GGATGCCCCG AAGGCGGCC TCTGAGCATT CCCTGGCCTC TGTGGCTGCC ACTTGGTTAT 660
 GTTGTGTGTG TCGGTGAGTG GTGTGCAGGC GCGGTCCTT ACCGCCATG TGTGCTGTGT 720
 GTGTCCAGGC ACGGTTCTT ACGCCCCATG TGTGCTGTGT GTTCTGCC TGTATATGTG 780
 50 GCTTCTCTG ATGCTGACAA GGTGGGGAAC AATCCTTCTT AATTTGGGT GGGACCAGAC 840
 TTTGTTCTCT TCCTCACCTG AAATTATGCT TCCTAAAT TTTAAATGAA CTCAAAGAAT 900
 55 GGGGTGGTGG GGGGCACCT GTGAGGTGGC CCCTGAGA TTTTCTTCTT TCCAGGGCAC 960
 ATCTGGAGTT CTTCTCCAGC TTACCCTAGG GTGACCAAT TTTTCTTCTT ACACCAGGT 1020
 60 GGCGCAGCTT TCTGTGTGAT GCAGATGTGT CCTGGTTTCT TTTTCTTCTT AGCTGCTGCT 1080

5 TGAGGCCATG GCTCCGTCCC CGGAGTTGGG GGTACCCGTT GCAGAGCCAG GGACATGATG 1140
 CAGGCGAAGT TCGGCCAAGTTGG ACTTTGATCC TTTGGGCAGA TGTCCCATTG 1200
 10 CTCCCTGGAG CCTGTCATGC CTGTTGGGGA TCAGGCAGCC TCCTGATGCC AGAACACCTC 1260
 AGGCAGAGCC CTA CTCTAGCT GTACCTGTCT GCCTGGACTG TCCCCTGTCC CCGCATCTCC 1320
 CCTGGGACCA GCTGGAGGGC CACATGCACA CACAGCCTAG CTGCCCCCAG GGAGCTCTGC 1380
 TGCCCTTGCT GGCCCTGCCC TTCCACAGG TGAGCAGGGC TCCTGTCCAC CAGCACACTC 1440
 AGTTCTCTTC CCTGCAGTGT TTTCATTTTA TTTTAGCCAA ACA TTTTGCC TGT TTTCTGT 1500
 15 TTCAAACATG ATAGTTGATA TGAGACTGAA ACCCCTGGGT TGTGGAGGGA AATTGGCTCA 1560
 GAGATGGACA ACCTGGCAAC TGTGAGTCCC TGCTTCCCGA CACCAGCCTC ATGGAATATG 1620
 CAACAACCTC TGTACCCAG TCCACGGTGT TCTGGCAGCA GGGACACCTG GGCCAATGGG 1680
 20 CCATCTGGAC CAAAGGTGGG GTGTGGGGCC CTGGATGGCA GCTCTGGCCC AGACATGAAT 1740
 ACCTCGTGTT CCTCCTCCCT CTATTACTGT TTCACCAGAG CTGTCTTAGC TCAAATCTGT 1800
 25 TGTGTTTCTG AGTCTAGGGT CTGTACACTT GTTTATAATA AATGCAATCG TTTGGAAAAA 1860
 AAAAAAAAAA AAACCTCGTAG GGGGGGCCCC TACCCAATGG GCYCMARAT AGTAGARWAC 1920
 RAAAAAYMCA ANTGCAACCA AAGAGGGGCC AGGGGANTTT TAAGAGGGCC CCCTTTTGGG 1980
 30 GGNATCCANT TTAGCCGGGG TTNTAAGGG AAGTTGCNTG GCGGGGGTTA GGGCCCSGTT 2040
 KYTWCTTCCA ACCAAGGGTT YTYGTGGTTA GGCCGGGTG GGCCCMATGG GCTGGGCTGG 2100
 35 GTAAAGTGGT GGGTMAYTGC MATTTGGGTAG GGTGCTGCTG GCATTCTCTG CTGAGGCGGC 2160
 ATGGTGTGGT AGCCCTGGTA GCTTGGTCCA GGTAGCTGG GCGGCACACT TGGAGGCTGA 2220
 GGATAAGGGG CATGCACCCA CAGTGGTGGA TGTGGTGGTG GTGACAACCG GACGTGGTCG 2280
 40 GCGGCACGTC TTGTAAAGGC AGCAGCAGGA GCAGGTGAAG CAGATGATGA TAGTGACGAC 2340
 AGACAGCACA AAGATGGTCC AGCCAACGGC CAAGGTCGCT CCGAACCCCA GTTTAGCCTC 2400
 45 TGCCTTCACC CAGAGGCCTC CTGTACCCTT GGGCTTTCCC CTC TTTCCCAT CGTGGGCCAC 2460
 TCACTCGTGC C 2471

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(2) INFORMATION FOR SEQ ID NO: 42:

55 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 2659 base pair.
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:

	GGCACCAGCT TTTCTCTAGA GTCTGAAAGA TGCTAGAAAG AAATAAAATT TAACTTACTT	60
5	AAGAGAATTA TGGATCTTTT ATTAATAAAA ATTAAC TTGA TGATTTGAAC TAACAGTTAT	120
	GATAATTCTG GTATTTTATAG CTTTTTTTAT TCCCCTGCAG AAAACCATAG GCAAAATTGC	180
	AACATGCTTG GAATTGCGAA GTGCAGCTTT ACAGTCCACA CAGTCTCAAG AAGAATTTAA	240
10	ACTGGAGGAC CTGAAGAAGC TAGAACCAAT CCTAAAGAAT ATTCTTACAT ATAATAAAGA	300
	ATTCCCATT GATGTTTACG CTGTCCCAT AAGAAGAATT TTGGCACCTG GTGAAGAAGA	360
15	GAATTTGGAA TTTGAAGAAG ATGAAGAAGA GGGTGGTGCT GGAGCAGGTC TCCTGATTCT	420
	TTCTTGCTAG AGTTCCCGGT ACTTTATTAC CAAGGTGACC ATCGGAACCA GGAATGACAT	480
	TACTCACTAT CAGAATTGAG AAAATTGGTT TGAAAGATGC TGGGCAGTGC ATCGATCCCT	540
20	ATATTACAGT TAGTGTAAG GATCTGAATG GCATAGACTT AACTCCTGTG CAAGATACTC	600
	CTGTGGCTTC AAGAAAAGAA GATACATATG TTCATTTTAA TGTGGACATT GAGCTCCAGA	660
25	AGCATGTTGA AAAATTAACC AAAGGTGCAG CTATCTTCTT TGAATTCAAA CACTACAAGC	720
	CTAAAAAAG GTTTACCAGC ACCAAGTGTT TTGCTTTCAT GGAGATGGAT GAAATTAAAC	780
	CTGGGCCAAT TGTAATAGAA CTATACAAGA ACCCCACTGA CTTTAAAAGA AAGAAATTGC	840
30	AATTATTGAC CAAGAAACCA CTTTATCTTC ATCTACATCA AACTTTGCAC AAGGAATGAT	900
	CCTGACATGA TGAACCTGGA ACTTCTGTGA ATTTTACCAC TCAGTAGAAA CCATCATAGC	960
35	TCTGTGTAGC ATATTCACCC TTCAACAGGC AGGAAGCAAG CCGTACCCAG ACCAGTAGGC	1020
	CGGACGGAGT CAAATGCAAA GCTGTACCAC AGAATTCAGA GTCCAGCACA TCACACTGAC	1080
	GTATAGGACT CCTTGGGATA CAGGTTTATT GTAGATTTTG AAACATGTTT TTACTTTTCT	1140
40	ATTAATTGTG CAATTAATAG TCTATTTTCT AATTTACCAC TACTCCTACC CTGCTTCCTG	1200
	GAACAATACT GTTGTGGGTA GGATGTGCTC ATCTTCAGAC TTAATACAGC AATAAGAATG	1260
45	TGCTAGAGTT TACACATCTG TTCACTTTGT CTCCAATATG CTCTTTTGAC TTAACGTCAA	1320
	GCTTTGGGTT GATGTGGGTA GGGTAGTGTC AAAC TGCTTT GAGAGGAATG GGACCAGTTC	1380
	TGCTGCCTAA GAAGGTCTGT CTGGATGTTT ATAGGCAGCA CCTCTGAAGT GGCCTAAATT	1440
50	CACCCTGATC TGATAGTTTT CCTGCTTAGA AAGTGTGCCT TGGCCAGATC AGTATCCCAC	1500
	ATGGGAGTGT TCCCTAGGTT GTAGCTGTGA TTGTTTCCAG ATGACCAGAT TGTTTTCTG	1560
55	AAAATGAGCA TATTTT TAGT CATGTCGATT AGCTGTTCTT CTACATCACA TTGTTACTCT	1620
	TTCTGATGAT GATTCTAGGG TTAACATTGG AACCATCTCA AAATAATTAC AAAGTTT TAG	1680
	ATGGGTTTAC AATGTCTTCT AAACAATGTA ATCTAAAAAT AATTGAGTCA GATGCTAACG	1740
60	AGATACTGCA GGCATAACTG CTGTTTTTCT GACAACTGAT TGTGAAACCT TAAAACCTGC	1800

	ATACCTCTTC TTACAGTGAG GAGTATGCAA AATCTGGAAA GATATTCTAT TTTTTTTATA	1860
5	TAGGTAGATA GGATCGCCAT TTATTTCCCTA TTTAGATATA CTGACATTCA TCCATATGAA	1920
	AATATGCAGG TCATTAGCTT ACTATAATTT ACTTTTGACT TAATGGGGCA TAAATAAAAC	1980
	TTTCATAGTA CACATGAGGT GGATATTTGA TACACAGAAC ATTTGCGGTG GGCTTTCTGT	2040
10	GGGTTAGATG TAAAGCCAC ATATTTTAAT ATTCACTATT TTAAATGAGC AATGCATGAG	2100
	GGGAATGCAG TGTCAGTACC TGGCCTATTT TTAAACTAGT GTAATCACCC TAGTCATACC	2160
15	ATTCAGTATG TTGCTTTTT AAAATAAGTA ACCACAATTA AGTTGTGTGA GCCCTTGCAC	2220
	TTCAAGAGAT CTAGTCTTTA CTTTCAGTTG TCTGTTAGGT CCATTCTGTT TACTAGACGG	2280
	ATGTTAATAA AACTATGCG AGCCTGGAAT GGAATTCTCC AGCCAAATTT TAGTCTTGTC	2340
20	CTCTCCATCT TGATTGGATT AATTCCAAAT TCTAAAATGA TTCAGTCCAC AATAGCTCTA	2400
	GGGGATGAAG AATTTGCCCT ACTTTGCCCA GTTCCTAAGA CTGTGAGTTG TCAAATCCCT	2460
25	AGACTGTAAG CTCTTCAAGG AGCAAGAGGC GCATTTTCTC CGTGTGATGT AATTTTCTA	2520
	AGGTGTTGG CAGCACTCTG TACCCTGTGG AGTACTCAGT ACCTTTTGTT TGATGTTGCT	2580
	GACAAGACCT GAAAAAAAT CCCTTAAAAA AAAAACCCAT TAAAGTGTAG CAAAACCGAA	2640
30	AWAAAAAAAA AAAAAAAAAA	2659
35	(2) INFORMATION FOR SEQ ID NO: 43:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1635 base pairs	
	(B) TYPE: nucleic acid	
40	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43:	
45	CGAGGAGGTC ATGAACAAGG AGGCGGGAGA GGTGGACGTG GTGGCTATGA CCATGGTGGC	60
	CGAGGGGGAG GAAGAGGAAA TAAGCATCAA GGAGGCTGGA CAGATGGAGG GAGTGGTGGG	120
50	GGAGGTGGCT ACCAAGATGG TGGTTATCGA GATTCAGGTT TCCAGCCAGG TGGCTATCAT	180
	GGTGGCCACA GCAGTGGTGG CTATCAAGGC GGAGGTTATG GTGGCTTCCA AACATCTTCT	240
	TCATATACAG GAAGTGGATA CCAGGGTGGT GGCTACCAGC AGGACAATAG ATACCAAGAT	300
55	GGCGGGCACC ATGGTGATCG TGGTGGTGGT CGTGGTGGG GAGTTGGTCG TGGAGGCCGA	360
	GGTGGTCGTG CAGGCCAGGG AGGAGGCTGG GGAGGAAGAG GGAGCCAGAA TTATCACCAA	420
60	GGGGGTCAAT TTGAACAGCA TTTCCAGCAT GGAGGTTATC AGTATAATCA TTCTGGATTT	480

	GGACAGGGAA GACATTACAC TAGTTGAGGC TACCGAACCT TACATTTTGC TAGAGCTCAA	540
	GTAATAGAAA CTTAGTTTCA GAATCCTGAA TTCAGCACCT ATTTTGAATT AATGTGAGAC	600
5	CACAGGTGGC AGGCAGATTC CTGCTTGGCA TAAGCATTTG TAGGTCTTCA TTCAATTCTG	660
	TTAGATTTTT TTATTGGACT TACATAATGC CGTTTATTTG AGAAACACAT AACATCTCTC	720
	CTTCTATGA AAAATTTTTT AAAAGGTGGT TAAAATTGCC TTTAATTGCC CAGTAGACTA	780
10	ATTCCACAGT CAGAACATGC AAACTTTTTT GAAGAAATTA CTTGAATAAG TAGTTTTCAT	840
	GTTTTCAATA TGCAGTTTTG AAAATGAGGA TTCACCTAGA CTTTTTTAGA TTTACTACYA	900
15	GGAAACCTTC CYCATATGAA TAACCATTTA TATGTGTTTT GCTTAAAGTA TTCCAATGCC	960
	TATTTTCCAA GCACAGTTCT GCCCCCGGT TGACTTTTAT GCCACGTGTG CTTTCATGATG	1020
	GAACTTTTAG GTCAGTTCCT ATTAAATGAG CTCCTTYTGCA GATAGCACAT TCAGTAGCCT	1080
20	TATTTTGTG ATGGAATACT GTATCATATG CTCAACTCTG AAAACCTTGA ACACGGCCAA	1140
	AATCCATAAA GATTATAAAA GCAAACCTAAG TTGTGAAGCT ATAGTACATG TAGGCATTTA	1200
25	GTTAAGTATA GCAATTCAAA CTGACCTGCA TCCATCCAAA ACAAATTCCT CCTTCAACCT	1260
	TATTTTTACT TGAAATTTGC TAGAAGAAAT AGCAAACCGA AATTTGTTTT ATGCATGAGT	1320
	TAATACCACT GGCTCAGCAA ATACAAGTTA GTTTGCTTTA AGCAGGTAAC TTTTTTTGTA	1380
30	ATGGAAGAAA TGCACCTACAA AGTTAAGACA GATTTTTGCT AAGTGCAGGA GGCCCTTTAT	1440
	TATTGCTGCA GAAAACAAAA GCCTGGCTGA GTTGATGTTT TACATTCTCC CTTACTGAAA	1500
35	TCTACATGAC ATGATGCTTC TTGCTGGGTT TTTGTACATG TAAACATTGT CAAGCTGTGA	1560
	AAGAAAATGG CTGGAGGTGT GCTTTGTGTG AAAGGTGAGC ACTGAAAGTA TCTGTTAAGT	1620
40	TCTCCNGAAA AAAAA	1635

45 (2) INFORMATION FOR SEQ ID NO: 44:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 780 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:

	AACATGGTCA TGTCTTTTAG TTTCATTATT TTCCTACTCC TTGTATGTCA AGAAATTACA	60
55	TTTTGCATGT CTTATGGAGA TGCTGTTAAT TGCTTCAGTG AGTGCTTTTC TAATCTGCAG	120
	ACCATTTACA TTTCTGTTT GCAGCATGCT GTGTGCAAAC ATTCAGTAAT TTGGAGTATT	180
60	CAATTATTTG TTAGGGCTCT TCCTATTTCC AAATGTGCTG AATTGCTAT TGATGGGATT	240

5 TTCAGATCTT TTCATGAGAA CTGGAAATGT AGCTGGGTGG CACCTACCTA GGTGCTACG 300
 TAGTGAGTAG ACTTCTCTT GGGTATAGTA AGCCTCAGAC AGCTTTCAC TTTATCTACT 360
 TTACTTGTGG AAATAAACA GTCATTTTGT TCTGAAAGAA TAAGATAGCT TTCTGTAGAG 420
 AAGGAATTCC TACCTCTAAA AGCTGCCTTG AGAACTCAGA ACTGGCAGTT TTCTGAGGTG 480
 10 ATTTTFAAAT TTCAGTATTA GGGAGAGTCC AGCATTTGCT GACACAGATT CTACATAACT 540
 AATGTATGAT AGCAAATGCA AACTATTAT AATGTGGTGT ATCTTGCGCA TACACAGGTT 600
 15 AGAACAAGTA GACTCTGGCA GCAGATCTCC AGAGACCCAA GTTTAGGTTT TCATAGTGTA 660
 TTTGAAGTAG TTATACTCCT GGCTTAAGTA GTTTAGTGCC TGGGAGAATC CATTACTGAA 720
 AAGCATTTAA CTAAAAAAA AAAAAAAAAA AAAACTGAAA AGGTAGTGAA TACAGAATAG 780

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(2) INFORMATION FOR SEQ ID NO: 45:

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- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 2378 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:

GCGAAGCAGC TGAAGCCGCC GCCGCGCAGA ATCCACGCTG GCTCCGTGCG CCATGGTCAC 60
 35 CCACAGCAAG TTTCCCGCCG CCGGGATGAG CCGCCCCCTG GACACCAGCC TGCGCCTCAA 120
 GACCTTCAGC TCCAAGAGCG AGTACCAGCT GGTGGTGAAC GCAGTGCACA AGTGCAGGAG 180
 AGCGGCTTCT ACTGGAGCGC AGTGACCGGC GCGAGGGCGA ACCTGCTGCT CAGTGCCGAG 240
 40 CCGCCCGCA CCTTCTGAT CCGCGACAGC TCGGGACCAG CGCCACTTCT TCACGCTCAG 300
 CGTCAAGACC CAGTCTGGA CCAAGAACCT GCGCATCCAG TGTGAGGGG GCAGCTTCTC 360
 45 TCTGCAGAGC GATCCCCGGA GCACGCAGCC CGTGSCCCGC TTCGACTGCG TGCTCAAGCT 420
 GGTGCACCAC TACATGCCGC CCCCTGGAGC CCCCTCCTTC CCTCGCCAC CTACTGAACC 480
 CTCCTCCGAG GTGCCCAGC AGCCGTCTGC CCAGCCACTC CCTGGGAGTC CCCCAGAAG 540
 50 AGCCTATTAC ATCTACTCCG GGGGCGAGAA GATCCCCCTG GTGTTGAGCC G3CCCCCTCTC 600
 CTCCAACGTG GCCACTCTTC AGCATCTCTG TCGGAAGACC GTCAACGCC ACCTGGACTC 660
 55 CTATGAGAAA GTCACCCAGC TGCCGGGGCC CATTCGGGAG TTCTTGAGCC AGTACGATGC 720
 CCCGCTTTAA GGGGTAAAGG GCGCAAAGG CATGGGTCGG GAGAGGGGAC GCAGGCCCTT 780
 CTCCTCCGTG GCACATGGCA CAAGCACAAG AAGCCAACCA GGAGAGAGTC CTGTAGCTCT 840
 60

GGGGGGAAAG AGGGCGGACA GGCCCTCCC TCTGCCCTCT CCCTGCAGAA TGTGGCAGGC 900
 GGACCTGGAA TGTGTTGGAG GGAAGGGGGA GTACCACCTG AGTCTCCAGC TTCTCCGAG 960
 5 GASCCAGCTG TCCTGGTGGG ACGATAGCAA CCACAAGTGG ATTCTCCTTC AATTCCTCAG 1020
 CTTCCCTCT GCCTCCAAAC AGGGGACACT TCGGAATGC TGAAC TAATG AGAACTGCCA 1080
 10 GGGAACTCTT AACTTTCCA ACGGAAGTGG TTTGCTCTTT GATTTGGTTT AAACCTGAGC 1140
 TGGTTGTGGA GCCTGGGAAA GGTGGAAGAG AGAGAGGTCC TGAGGGCCCC AGGGCTGCGG 1200
 GCTGGCGAAG GAAATGGTCA CACCCCCCGC CCACCCAGG CGAGGATCCT GGTGACATGC 1260
 15 TCCTCTCCCT GGCTCCGGGG AGAAGGGCTT GGGGTGACCT GAAAGGGAAC CATCCTGGTG 1320
 CCCCACATCC TCTCTCCGG GACAGTCACC GAAAACACAG GTTCCAAAGT CTACCTGGTG 1380
 CCTGAGAGCC CAGGGCCCTT CCTCCGTTTT AAGGGGGAAG CAACATTTGG CACGAGATGG 1440
 20 GCTGGTCAGC TGGTCTCCTT TTCCTACTCA TACTATACCT TCCTGTACCT GGTGGATGG 1500
 AGCGGGAGGA TGGAGAGACG GGACATCTTT CACCTCAGGC TCCTGGTAGA GAATACAGGG 1560
 25 GATTCTACTC TGTGCCTCCT GACTATGTCT GGCTAAGAGA TTCGCCTTAA ATGCTCCCTG 1620
 TCCCATGGAG AGGGACCCAG CATAGGAAAG CCACATACTC AGCCTGGATG GGTGGAGAGG 1680
 CTGAGGGACT CACTGGAGGG CACCAAGCCA GCCACAGCC AGGGAAGTGG GGAGGGGGGC 1740
 30 GGAAACCCAT GCCTCCAGC TGAGCACTGG GAATGTCAGC CCAGTAAGTA TTGGCCAGTC 1800
 AGGCGCCTCG TGGTCAGAGC AGAGCCACCA GTTCCACTG CCCCAGCCC TGCACAGCCC 1860
 35 TCCCTCCTGC CTGGGTGGGG GAGGCTGGAG GTCATTGGAG AGGCTGGACT GCTGCCACCC 1920
 CGGGTGCTCC CGCTCTGCCA TAGCACTGAT CAGTGACAAT TTACAGGAAT GTAGCAGCGA 1980
 TGAATTACC TGAACAGTT TTTGTTTTT GTTTTTGTTT TTGTTTTTGT GGGGGGGGGC 2040
 40 AACTAAACAA ACACAAAGTA TTCTGTGTCA GGTATTGGGC TGGACAGGGC AGTTGTGTGT 2100
 TGGGGTGGTT TTTTCTCTA TTTTGTGTG TGTTCCTTGT TTTTAAATAA TGTTTACAAT 2160
 45 CTGCCTCAAT CACTCTGTCT TTTATAAAGA TTCCACTCCA GTCCTCTCTC CTCCCCCTA 2220
 CTCAGGCCCT TGAGGCTATT AGGAGATGCT TGAAGAACTC AACAAAATCC CAATCCAAGT 2280
 CAACTTTGC ACATATTTAT ATTTATATTC AGAAAAGAAA CATTTAGTA ATTTATAATA 2340
 50 AAGAGCACTA TTTTAAATG AAAAAAAAAA AAAAAAAA 2378

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(2) INFORMATION FOR SEQ ID NO: 46:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1772 base pairs

(B) TYPE: nucleic acid

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(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:

5	TCGACCCACG CGTCCGGGAG GATCCCCAGC CGGGTCCCAA GCCTGTGCCT GAGCCTGAGC	60
	CTGAGCCTGA GCCGAGCCGG GAGCCGGTCG CGGGGGCTCC GGGCTGTGGG ACCGCTGGGC	120
10	CCCCAGCGAT GGCAGCCCTG TGGGGAGGCC TTCTTCGGCT TGGCTCCTTG CTCAGCCTGT	180
	CGTGCCCTGGC GCTTTCCTGT CTGCTGCTGG CGCACTGTCA GACGCCGCCA AGAATTTCTGA	240
15	GGATGTCTAGA TGTAAATGTA TCTGCCCTCC CTATAAAGAA AAATTCTGGG CATATTTATA	300
	ATAAGAACAT ATCTCAGAAA GATTGTGATT GCCTTCATGT TGTGGAGCCC ATGCCTGTGC	360
	GGGGGCCTGA TGTAGAAGCA TACTGTCTAC GCTGTGAATG CAAATATGAA GAAAGAAGCT	420
20	CTGTCACAAT CAAGGTTACC ATTATAATTT ATCTCTCCAT TTTGGGCCTT CTACTTCTGT	480
	ACATGGTATA TCTTACTCTG GTTGAGCCCA TACTGAAGAG GCGCCTCTTT GGACATGCAC	540
	AGTTGATACA GAGTGATGAT GATATTGGGG ATCACCAGCC TTTTGCAAAT GCACACGATG	600
25	TGCTAGCCCG CTCCCGCAGT CGAGCCAACG TGCTGAACAA GGTAGAATAT GGCACAGCAG	660
	CGCTGGAAGC TTCAAGTCCA AGAGCAGCGA AAAGTCTGTC TTTGACCGGC ATGTTGTCTT	720
30	CAGCTAATTG GGGAAATTGAA TTCAAGGTGA CTAGAAAGAA ACAGGCAGAC AACTGGAAAG	780
	GAACTGACTG GGTMTTGCTG GGTTCATTT TAATACCTTG TTGATTCAC CAACTGTTGC	840
	TGGAAGATTC AAAACTGGAA GKAAAACTT GCTTGATTTT TTTTCTTGT TAACGTAATA	900
35	ATAGAGACAT TTTTAAAAGC ACACAGCTCA AAGTCAGCCA ATAAGTCTTT TCCTATTTGT	960
	GACTTTTACT AATAAAAATA AATCTGCCTG TAAAATAAAT TAAAAAATCC TTTACCTGGA	1020
40	ACAAGCACTC TCTTTTTCAC CACATAGTTT TAACTTGACT TTCCAAGATA ATTTTCAGGG	1080
	TTTTTGTGTG TGTGTTTTTT TGTTTGTTTG TTTTGGTGGG AGAGGGGAGG GATGCCTGGG	1140
	AAGTGGTTAA CAACTTTTTT CAAGTCACTT TACTAAACAA ACTTTTGTAA ATAGACCTTA	1200
45	CCTTCTATTT TCGAGTTTCA TTTATATTTT GCAGTGTAGC CAGCCTCATC AAAGAGCTGA	1260
	CTTACTCATT TGACTTTTGC ACTGACTGTA TTATCTGGGT ATCTGCTGTG TCTGCACTTC	1320
50	ATGGTAAACG GGATCTAAAA TGCCTGGTGG CTTTTCACAA AAAGCAGATT TTCTTCATGT	1380
	ACTGTGATGT CTGATGCAAT GCATCCTAGA ACAAACGGC CATTTGCTAG TTTACTCTAA	1440
	AGACTAAACA TAGTCTTGGT GTGTGTGGTC TTACTCATCT TCTAGTACCT TTAAGGACAA	1500
55	ATCCTAAGGA CTTGGACACT TGCAATAAAG AAATTTTATP TTAAACCCAA GCCTCCCTGG	1560
	ATTGATAATA TATACACATT TGTCAGCATT TCCGGTCGTG GTGAGAGCCA GCTGTTTGAG	1620
60	CTCCAATGTG TGCAGCTTTG AACTAGGGCT GGGTTGTGG GTGCCCTCTT TGAAAGGTCT	1680

AACCATTATT GGATAACTGG CTTTTTTTCT TCCTCTTTGG AATGTAACAA TAAAAATAAT 1740
 TTTTGAAACA TCAAAAAAAA AAAAAAAA AA 1772
 5

 (2) INFORMATION FOR SEQ ID NO: 47:
 10
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1107 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 15 (D) TOPOLOGY: linear

 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:

 CGGGCGAGAA GGGCAGACGG GACATGCAGC CTCTTCCGCC TGAGCCCCGG AAGTGATGTG 60
 20 GCTGCGGCAT CGCGGCCTCG CTATGTCTGC CATTTTCAAT TTTCAGAGTC TATTGACTGT 120
 AATCTTGCTG CTTATATGTA CCTGTGCTTA TATTCGATCC TTGGCACCCA GCCTCCTGGA 180
 25 CAGAAATAAA ACTGGATTGT TGGGTATATT TTGGAAGTGT GCCAGAATTG GTGAACGGAA 240
 GAGTCCTTAT GTTGCACTAT GCTGTATAGT AATGGCCTTC AGCATCCTCT TCATACAGTA 300
 GCTGGGGAAA ATGCCAGAAT GTAGTTGCCA TCAGATTTGA TTGTGAACAA GGACTGACTG 360
 30 CAGAAAATAA TGGAAAGGAT GTTTAACTCT TTTATCTCCG AACATTGAAT GAGATAAATT 420
 TCCAGATGCT GTTCTCTATT TTAATGTTAT TGGACCAATG TTCTGTATAA ACAATTAAGA 480
 35 TGTAACCATT TAATAGTCTG TAACAATCAA CCTCAGTACT GTCACTACAA TATTACATTG 540
 TGCAATGTT ATTCTGTTGT ATCAGATACA AAATTTTAGT GAGGTATCTC TAAGGCACAT 600
 AGTAGAAAAC AAAATTGGTT AATTACTCAA GTTCCTTTCA CTGTGATTIG GAAATGATTT 660
 40 AATCTTTATA GAATGAGAAC CTTTTTGGGA CTAGCTTTTT TATTAAAATG GCTCAATTTG 720
 TGTGATAAG GATTGCATTA ATATTTAATA GTGCTTGCTT TCCTCTGGG CACACCATTT 780
 45 TGATCATTAA CCAGAGTACC TCTACTCTTA GCAAACCTTA GTTTATGACA AGTATTTAAA 840
 ATATTTAAAA CAAGCTTATG CAGTTCTTAA GGACGAAGGT AAATGAGATG TAACTTAAAA 900
 ATAGTATTGG GAAAATGTTG ATAGTTAACA TTAGTGGAAT TAGACTAGCC AAATGACATA 960
 50 GTAGGCTCTG AAACATCTTG TCAAGTATAT GTATTTTGTG CATGAATTTT TGCTGGAAAG 1020
 CTGTCTTTCT CTGAAAAACA CAACCTTCTT AGAATGAAAA GAACAATTAT AAAATAAAAA 1080
 55 AAAAATTTAA AAAAACTGG GCGGGGG 1107

60 (2) INFORMATION FOR SEQ ID NO: 48:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 805 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:

10 TGCAGAAGAG ATGGAGTTGC TGTTGGAAAA CTACTACCGA TTGGCTGACG ATCTCTCCAA 60
TGCAGCTCGT GAGCTTAGGG TGCTGATTGA TGATTACAAA AGTATTATTT TCATTAATCT 120
15 GGACAGCCAC CGAAACGTGA TGATGAGGTT GAATCTACAG CTGACCATGG GAACCTTCTC 180
TCTTTCGCTC TTTGGACTAA TGGGAGTTGC TTTTGAATG AATTGGAAT CTTCCTTGA 240
AGAGGACCAT AGAATTTTTT GGCTGATTAC AGGAATTATG TTCATGGGAA GTGGCCTCAT 300
20 CTGGAGGCGC CTGCTTTCAT TCCTTGACG ACAGCTAGAA GCTCCATTGC CTCCTATGGT 360
ATGAAGGATA TGGTTCACGG CGGTATTGTG GAAGGGTTAT GATCATGGGC CCTAAAGTCA 420
GAGCGCCTGG GATTAAGTTG TCACAGGCAC TATGGCCCTT GCGAGTTGCT TTCTCAAAC 480
25 TCCTTCAGTT TCCCTATCTG TCAGTTAAGT CGGTATTACC TGCTTCATAG GGTATGGGA 540
AGAATTAAAC AATATGTGTA AAGCACTTAC TAGCACACTG CCTAACACAA TAAGTTAGAA 600
30 ATATAATTTG TGTAGAACTC TGACAACATA CATTTAAACA GATGTTAGTA ATTCTGGTAT 660
AAGGTTTGTC ATAACCAAAT GGAAATGTAG GAAACATTTA TAATGTTCTT AAAAGATAGR 720
AAATTCACCT CCATTTTCTT TGTACTTGAA GATGGCACCA CTGGAATAAA TACTTAAGAC 780
35 ACTGAAAAAA AAAAAAAAAA AACTC 805

40

(2) INFORMATION FOR SEQ ID NO: 49:

(i) SEQUENCE CHARACTERISTICS:

- 45 (A) LENGTH: 1408 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:

50 TCATTATTTA TTCATGTGGC TGAAAGAGTA TATTAATTAT GTTTAGATTT TTGAAAAAG 60
TCTGAACAAA AAAAGGACCT ATACAGTGCT CAACTATAT TTTTAAAAAT ACTATTTTAT 120
55 TTTTACTCAC ATATGAAAAA AATGGCTGTA CTATCATGTT TAATATAGCA CTAACATTGG 180
AAACAGAATA ACGAATTGTA TTAAATTTT ATGAAGAACA CACAAACATT AAAACACTGA 240
60 TTGGTTACAG AAAGCAGAGT TTGAGGAAAA AACATTAGCT ATAATTTTCA TTTTCATTAA 300

195

AGAGCAGCAC CCTCTGAGAA TAATCAAACCT GATTAGTAAT ATTCATCTAT ACTGCAAAAT 360
 AATATGTACA AAGGAAAGTT AGTGATTGTA CTGATTTTAT TACTTTTACC AAGCCATTTT 420
 5 ATGTTTCCTCA CTCAATGCAA AGAAATAAAA CATAATCTGA AGAAAAATAT GTCCTTATTA 480
 TTATTACAAA TAAAAAGTTG GCTTTATTCT GCAAGCCTGG GCATATTGTA CAATTGGCAG 540
 CACTTAACGG CTCAAGTGA TCAATGTACC AGTTTGATTC TGATCCACTG AATAGAATCT 600
 10 CTCATCCATA TCTGGTGACC AGACTAACTC CATGGGAGCT GTGATAGACT GAACCATTTT 660
 TGTGGTATCC CTAGATCTCA CTAAATAAGA AAGACCCTAC ACCAGAAAAT ATAGCAACTG 720
 15 ATCTATCTAT AAATTACATC TATATGCTAG CTCTTTAGTA TAAGTTGGAA AAAGGGGCCC 780
 TTTCTTGAGC ACATGGATAA AAGTATTATT GTAGTCTAAA GATTGCTGGA TTGATATTGT 840
 GTTGTATATA TGAAGATAAG GTACACACTG AAACCACTGT CAGATTAAGA AACTTCCACA 900
 20 ACTTGTCTCA GTTCTTCAAA CAATGGAGCA AGTTCCTTTT CTAGGCTGAC AATTAGTCCT 960
 GTATTGGCAC TGCTGCTGGC TATGAAACTC ACCACCAAAG GTAAACGATT AAATTGAACC 1020
 25 ACCTGGTAGG TGTTATAGTA ACAGATGATA CTTTTATTTT TGGAAAGTCC AAGTTTGCTT 1080
 CCTTGGTCTG TTGCAAGGGC AAAAGTGGAT AAGAAACCAG GTCGCAAAGC ATGCTCTGGA 1140
 GCATTGTCAT TTGCCACTTT AATAACAGGT ACTCCATCTC TATCTGACAC AACAATGGCA 1200
 30 TGGAGCCCTT CAACACTTGG TAACTTTTTA TACAAGAATC GCTTTAGGTC ATCCGCCATG 1260
 ATGAACCCCC TTCTCTCGCA GGATCAATCT CCACGCCTGG GGTTCCTGGG CTGCCTGGTT 1320
 35 CTCTCCGCTG TCACTTCAGG GACAGCTTTA AAGACAGGTT CCTCCTCAAG CCACCGTCAC 1380
 ATGATTCATG ACCTCGTCTG CGCTCCAG 1408

40

(2) INFORMATION FOR SEQ ID NO: 50:

45 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1813 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:

CATGGTGGGG CACGAGATGG CCTCTRACTC TTCWAACACTT TAAAGGCTAT TCTCAAACAT 60
 GGGAAATCCA ATGAACACCA CACAGTTAGG GAAATCACTT TTTTATTTT AGGTGGAGCA 120
 55 GGAAGAAAGC AAATTGGCAA ATATTTCCCA AGACCAGTTT TTTTAAAAAT ATGCAGATGG 180
 TGACACGTTT CTTCATATTG CTGTTGCCCA AGGGAGAAAT TTTTCTTCT ATGTTCTTGC 240
 60 AAGAAAGATG AATGCACTTC ACATGCTGGA TATTAAAGA TAAATATGAC AGAGTGCCTT 300

	TCAGGTGGCA GTGGCTGCCA ATCAGCATCT CATGTGCGAG GATCTGGTGA ACATCGGGGC	360
5	ACAGGTGAAC ACCACAGACT GCTGGGGAAG AACACCTCTG CATGTGTGTG CTGAGAAGGG	420
	CCACTCCCAG GTGCTTCAGG CGATTTCAGAA GGGAGCAGTG GGAAGTAATC AGTTTGTGGA	480
	TCTTGAGGCA ACTAACTATG ATGGCCTGAC TCCCCTTCAC TGTGCAGTCA TAGCCCACAA	540
10	TGCTGTGGTC CATGAACTCC AGAGAAATCA ACAGCCTCAT TCACCTGAAG TTCAGGAGCT	600
	TTTACTGAAG AATAAGAGTC TGGTTGATAC CATTAAAGTGC CTAATTCAAA TGGGAGCAGC	660
15	GGTGAAGCG AAGGATCGCA AAAGTGCCCG CACAGCCCTG CATTTTGGCAG CTGAAGAAGC	720
	AAATCTGGAA CTCATTCGCC TCTTTTTTGA GCTGCCCAGT TGCCTGTCTT TTGTGAATGC	780
	AAAGGCTTAC AATGGCAACA CTGCCCTCCA TGTGCTGCC AGCTTGCACT ATCGGTTGAC	840
20	ACAATTAGAT GCTGTCCGCC TGTTGATGAG GAAGGGAGCA GACCCAAGTA CTCGGAACCT	900
	GGAGAACGAA CAGCCAGTGC ATTTGGTTCC CGATGGCCCT GTGGGAGAAC AGATCCGACG	960
25	TATCCTGAAG GGAAAGTCCA TTCAGCAGAG AGCTCCACCG TATTAGCTCC ATTAGCTTGG	1020
	AGCCTGGCTA GCAACACTCA CTGTCAGTTA GGCAGTCCCTG ATGTATCTGT ACATAGACCA	1080
	TTTGCCTTAT ATTGGCAAAT GTAAGTTGTT TCTATGAAAC AAACATATTT AGTTCACTAT	1140
30	TATATAGTGG GTTATATTAA AAGAAAAGAA RAAAAATATC TAATTWCTCT TGGCAGATTT	1200
	GCATATTTCA TACCAGGTA TCTGGATCTA GACATCTGAA TTTGATCTCA ATGGTAACAT	1260
35	TGCCTTCAAT TAACAGTAGC TTTTGAGTAG GAAAGGACTT TGATTGTGG CACAAAACAT	1320
	TATTAATATA GCTATTGACA GTTTCAAAGC AGGTAAATTG TAAATGTTTC TTTAAGAAAA	1380
	AGCATGTGAA AGGAAAAAGG TAAATACAGC ATTGAGGCTT CATTTGGCCT TAGTCCCTGG	1440
40	GAGTTACTGG CGTTGGACAG GCTTCAGTCA TTGGACTAGA TGAAAGGTGT CCATGGTTAG	1500
	AATTTGATCT TTGCAAACTG TATATAATTG TTATTTTGT CCTTAAAAAT ATTGTACATA	1560
45	CTTGGTTGTT AACATGGTCA TATTTGAAAT GTATAAGTCC ATAAAAATAGA AAAGAACAAG	1620
	TGAATTGTTG CTATTTAAAA AAATTTTACA ATTCTACTA AAGAGTTTTT ATTGTGTAAT	1680
	CACTAAGTCT TTGTAGATAA AGCAGATGGG GAGTTACGGA ATTGTTCCCT TACTGGCTGA	1740
50	AAGATATATT CGAATTGTAA AGATGCTTTT YCTCATGCAT TAAAAATATA CATTATTTGT	1800
	AGGGAATTGC ATG	1813

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(2) INFORMATION FOR SEQ ID NO: 51:

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 2070 base pairs

RNSMCHD-WO Q23QAAA2 1

GTCACCCATC TTCCAGTAAG CAGTTTATTC TGAGCCCCGG GGGTAGACAG TCCTCAGTGA 1680
 GGGGTTTTGG GGAGTTTGGG GTCAAGAGAG CATAGGTAGG TTCCACAGTT ACTCTTCCCA 1740
 5 CAAGTTCCCT TAAGTCTTGC CCTAGCTGTG CTCTGCCACC TTCCAGACTC ACTCCCCTCT 1800
 GCAAATACCT GCATTTCTTA CCCTGGTGAG AAAAGCACAA GCGGTGTAGG CTCCAATGCT 1860
 10 GCTTTCCCAG GAGGGTGAAG ATGGTGCTGT GCTGAGGAAA GGGGATGCAG AGCCCTGCCC 1920
 AGCACCACCA CCTCCTATGC TCCTGGATCC CTAGGCTCTG TTCCATGAGC CTGTTGCAGG 1980
 TTTTGGTACT TTAGAAATGT AACTTTTTCG TCTTATAATT TTATTTTATT AAATTAAATT 2040
 15 ACTGCAAAAA AAAAAAAAAA AAAAAAAAAA 2070

20 (2) INFORMATION FOR SEQ ID NO: 52:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1426 base pairs
 (B) TYPE: nucleic acid
 25 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:

30 CCCTCACTAA AGGGAACAAA AGCTGGAGCT CCACCGCGGT GCGGCCGCT CTAGAACTAG 60
 TGGATCCCCC GGGCTGCAGG AATTCGGCAC ACGGATCGGC GTCCGCAGCG GCGGCTGCT 120
 GAGCTGCCTT GAGGTGCAGT GTTGGGGATC CAGAGCCATG TCGACCTGC TACTACTGGG 180
 35 CCTGATTGGG GGCCTGACTC TCTTACTGCT GCTGACGCTG CTGGCCTTTG CCGGGTACTC 240
 AGGGCTACTG GCTGGGGTGG AAGTGAGTGC TGGGTCACCC CCCATCCGCA ACGTCACTGT 300
 40 GGCCTACAAG TTCCACATGG GGCTCTATGG TGAGACTGGG CGGCTTTTCA CTGAGAGCTG 360
 CAGCATCTCT CCCAAGCTCC GCTCCATCGC TGTCTACTAT GACAACCCCC ACATGGTGCC 420
 CCCTGATAAG TGCCGATGTG CCGTGGGCAG CATCTGAGT GAAGGTGAGG AATCGCCCTC 480
 45 CCCTGAGCTC ATCGACCTCT ACCAGAAATT TGGCTTCAAG GTGTTCCTCT TCCCGGAACC 540
 CAGCCATGTG GTGACAGCCA CTTTCCCTT AACACCACCA TTCTGTCCCA TCTGGCTGGG 600
 50 CTACCCGCCG TGTCCATCCT GCCTTGGACA CCTACATCAA GGAGCGGAAG CTGTGTGCCT 660
 ATCTCGGCT GGGATCTAC CAGGAAGACC AGAATCCATT TCATGTGCCC ACTGGCACGG 720
 CCAGGGAGAC TTCTATGTGC CTGAGATGAA GGAGACAGAG TGGAAATGGC GGGGGCTTGT 780
 55 GGAGGCCATT GACACCCAGG TGGATGGCAC AGGAGCTGAC AATATGACTG ACACGAGTTC 840
 TGTAAGCTTG GAAGTGAGCC CTGGCAGCCG GGAGACTTCA GGTGCCACAC TGTCACCTGG 900
 60 GCGGAGCAGC CGTGGCTGGG ATGACGGTGA CACCCGCAGC GAGCACAGCT AACAGCGAGT 960

5 CAGGTGCCAG CGGCTCCTCT TTGAGGAGC TGGACTTTGG AGGGCGAGGG GCCCTTAAGG 1020
 GGAGTCACGG CTGGACCCTG GGAAGTGGAG CCCTGGGGGA CTACCAAGTG GCTCTGGGAG 1080
 CCCACTGCCC CTGAGAAGGG CAAGGAGTAA CCCATGGCCT GCACCCTCCT GCAGTGCAGT 1140
 TGCTGAGGAA CTGAGCAGAC TCTCCAGCAG ACTCTCCAGC CCTCTTCCTC CTTCTCTGG 1200
 10 GGAHAGAGGG GTTCCTGAGG GACCTGACTT CCCCTGCTCC AGGCCTCTTG CTAAGCCTTC 1260
 TCCTCACTGC CCTTTAGGCT CCCAGGGCCA GAGGAGCCAG GGACTATTTT CTGCACCAGC 1320
 CCCCAGGGCT GCCGCCCTG TTGTGTCTTT TTTTCAGACT CACAGTGGAG CTTCCAGGAC 1380
 15 CCAGAATAAA GCCAATGATT TACTTGTTAA AAAAAAAAAA AAAAAA 1426

20

(2) INFORMATION FOR SEQ ID NO: 53:

25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1720 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:

GGCACGAGTG CGGCCCCAGC CTCTCCTCAC GCTCGCGCAG TCTCCGCCGC AGTCTCAGCT 60
 GCAGCTGCAG GACTGAGCCG TGCACCCGGA GGAGACCCCC GGAGGAGGCG ACAAACCTTCG 120
 35 CAGTGCCGCG ACCCAACCCC AGCCCTGGGT AGCCTGCAGC ATGGCCCAGC TGTTCCTGCC 180
 CCTGCTGGCA GCCCTGGTCC TGGCCCAGGC TCCTGCAGCT TTAGCAGATG TTCTGGAAGG 240
 40 AGACAGCTCA GAGGACCGCG CTTTTCGCGT GCGCATCGCG GGCGACGCGC CACTGCAGGG 300
 CGTGCTCGGC GGCGCCCTCA CCATCCCTTG CCACGTCCAC TACCTGCGGC CACCGCCGAG 360
 CCGCCGGGCT GTGCTGGGCT CTCGCGGGT CAAGTGGACT TTCTGTCCC GGGGCCGGGA 420
 45 GGCAGAAGTG CTGGTGGCGC GGGGAGTGCG CGTCAAGGTG AACGAGGCCT ACCGTTCCG 480
 CGTGGCACTG CCTGCGTACC CAGCGTCGCT CACCGACGTC TCCCCTGGCG CTGAGCGAGC 540
 TGCGCCCCAA CGACTCAGGT ATCTATCGCT GTGAGGTCCA GCACGGCATC GATGACAGCA 600
 50 GCGACGCTGT GGAGGTCAAG GTCAAAGGTA TCCCATCCAG ACCCCACGAG AGGCCTGTTA 660
 CGGAGACATG GATGGCTTCC CCGGGGTCCG GAACTATGGT GTGGTGGACC CGGATGACCT 720
 55 CTATGATGTG TACTGTTATG CTGAAGACCT AAATGGAGAA CTGTTCTCTG GTGACCCTCC 780
 AGAGAAGCTG ACATTGGAGG AAGCACGGGC GTACTGCCAG GAGCGGGGTG CAGAGATTGC 840
 CACCACGGGC CAACTGTATG CAGCCTGGGA TGGTGGCCTG GACCACTGCA GCCCAGGGTG 900
 60

	GCTAGCTGAT	GGCAGTGTGC	GCTACCCCAT	CGTCACACCC	AGCCAGCGCT	GTGGTGGGGG	960
	TTGCCTGGT	GTCAAGACTC	TCTTCTCTTT	CCCCAACCAG	ACTGGCTTCC	CCAATAAGCA	1020
5	CAGCCGCTTC	AACGTCTACT	GCTTCCGAGA	CTCGGCCCCAG	CTTCTGCCAT	CCCTGAGGCC	1080
	TCCAACCCAG	CCTCCAACCC	AGCTTTGATG	GACTAGAGGC	TATCGTCACA	GTGACAGAGA	1140
10	CCCTGGAGGA	ACTGCAGCTG	CCTCAGGAAG	CCACAGAGAG	TGAATCCCGT	GGGGCCATCT	1200
	ACTCCATCCC	CATCATGGAG	GACGGAGGAG	GTGGAAGCTC	CACTCCAGAA	GACCCAGCAG	1260
	AGGCCCTTAG	GACGCTCCTA	GAATTTGAAA	CACAATCCAT	GGTACCGCCC	ACGGGGTTCT	1320
15	CAGAAGAGGA	AGGTAAGGCA	TTGGAGGAAG	AAGAGAAATA	TGAAGATGAA	GAAGAGAAAG	1380
	AGGAGGAAGA	AGAAGAGGAG	GAGGTGGAGG	ATGAGGCTCT	GTGGGCATGG	CCCAGCGAGC	1440
20	TCAGCAGCCC	GGGCCCTGAG	GCCTCTCTCC	CCACTGAGCC	AGCAGCCCAG	GAGGAGTCAC	1500
	TCTCCCAGGC	GCCAGCAAGG	GCAGTCTTGC	AGCCTGGTGC	ATCACCCTT	CCTGATGGAG	1560
	AGTCAGAACC	TTCCAGGCCT	CCAAGGGTCC	ATGGACCACC	TACTGAGACT	CTGCCCCACTC	1620
25	CCAGGGAGAG	GAACCTAGCA	TCCCCATCAC	CTTCCACTCT	GGTTGAGGCA	AGAGAGGTGG	1680
	GGGAGGC AAC	TGGTGGTCTT	GAGCTATCTG	GGTCCCTCGA			1720

(2) INFORMATION FOR SEQ ID NO: 54:

(i) SEQUENCE CHARACTERISTICS:

35 (1) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1117 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:

	GGCACGAGGC CAAACTTCGG GCGGCTGAGG CGGCGGCCGA GGAGCGGCGG ACTCCGGGCG	60
45	CGGGGAGTCG AGGCATTTGC GCCTGGGCTT CGGAGCGTAC CCAGGGCCTG AGCCTTTGAA	120
	GCAGGAGGAG GGGAGGAGAG AGTGGGGCTC CTCTATCGGG ACCCCCTCCC CATGTGGATC	180
	TGCCCAGGCG GCGGCGGCGG AGGAGGCGAC CGAGAAGATG CCCGCCCTGC GCCCCGCTCT	240
50	GCTGTGGGCG CTGCTGGCGC TCTGGCTGTG CTGCGCGACC CCCGCGCATG CATTGCAGTG	300
	TCGAGATGGC TATGAACCCT GTGTAAATGA AGGAATGTGT GTTACCTACC ACAATGGCAC	360
55	AGGATACTGC AAAGGTCCAG AAGGCTTCTT GGGGGAATAT TGTCAACATC GAGACCCCTG	420
	TGAGAAGAAC CGCTGCCAGA ATGGTGGGAC TTGTGTGGCC CAGGCCATGC TGGGGAAAGC	480
	CACGTGCCGA TGTGCCTCAG GGTTTACAGG AGAGGACTGC CAGTACTCGA CATCTCATCC	540
60	ATGCTTTGTG TCTCGACCTT GCCTGAATGG CGGCACATGC CATATGCTCA GCCGGGATAC	600

201

CTATGAGTGC ACCTGTCAAG TCGGGTTTAC AGGTAAGGAG TGCCAATGGA CCGATGCCTG 660
 CCTGTCTCAT CCCTGTGCAA ATGGAAGTAC CTGTACCACT GTGGCCAACC ATTTCTTGCA 720
 5 AATGCCTCAC AGGCTTCACA GGGCAGAAGT GTGAGACTGA TGTCAATGAG TGTGACATTC 780
 CAGGACACTG CCAGCATGGT GGCACCTGCC TCAACCTGCC TGGTTCCTAC CAGTGCCAGT 840
 10 GCCTTCAGGG CTTCACAGGC CAGTACTGTG ACAGCCTGTA TGTGCCCTGT GCACCCTCGC 900
 CTTGTGTCAA TGGAGGCACC TGTCGGCAGA CTGGTGACTT CACTTTTGAG TGCAACTGCC 960
 TTCCAGAAAC AGTGAGAAGA GGAACAGAGC TCTGGGAAAG AGACAGGGAA GTCTGGAATG 1020
 15 GAAAAGAACA CGATGAGAAT TAGACACTGG AAAATATGTA TGTGTGGTTA ATAAAGTGCT 1080
 TTAAACTGAA AAAAAAAAAA AAAAAAAAAA AAAAAA 1117
 20

(2) INFORMATION FOR SEQ ID NO: 55:

25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1903 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:

GGCACGAGCT CGGAGAGGCG GCGCCCCTGA GTAGGCCAGG AGCCTCTCTT GCAACTTCTG 60
 35 CCACCGCGGG CCACCGCGGC CGCCTGATCC CGCAGAGGAA GGTGCGGGCC GTGGAGCGAT 120
 GACCCGCGGC GGTCCGGGCG GGCGCCCGGG GCTGCCACAG CCGCCGCCGC TTCTGCTGCT 180
 GCTGCTGCTG CCGCTGTTGT TAGTCACCGC GGAGCCCGCG AAACCTGCAG GAGTCTACTA 240
 40 TGCAACTGCA TACTGGATGC CTGCTGAAAA GACAGTACAA GTCAAAAATG TAATGGACAA 300
 GAATGGGGAC GCCTATGGCT TTTACAATAA CTCTGTGAAA ACCACAGGCT GGGGCATCCT 360
 45 GGAGATCAGA GCTGGCTATG GCTCTCAAAC CCTGAGCAAT GAGATCATCA TGTTTGTGGC 420
 TGGCTTTTTG GAGGGTTACC TCATTGCCCC ACACATGAAT GACCACTACA CAAACCTCTA 480
 CCCACAGCTG ATCACGAAAC CTTCATCAT GGATAAACTG CAGGATTTTA TGGAGAAGCA 540
 50 AGATAAGGTG GACCCGAAA AATATCAAAG AATACAAGAC TGATTCATTT TGGAGACATA 600
 CAGGCTATGT GATGGCACA ATAGATGGCC TCTATGTAGG AGCAAAGAAG AGGGCTATAT 660
 55 TAGAAGGGAC AAAGCCAATG ACCCTGTTCC AGATTCAGTT CCTGATAAT GTTGGAGATC 720
 TATTGGATCT GATTCCCTCA CTCTCTCCCA CAAAAACCG CAGCCTAAAG GTTTTAAAGA 780
 GATGGGACAT GGGACATTGC TCCGCTCTTA TCAAGGTTCT TCTTGTGTTT GAGAACATCC 840
 60

TTTTGTCTCA CTCAAGCTGG TACACGTATG CAGCCATGCT CAGGATATAT AAACACTGGG 900
 ACTTCAACAT CATAGATAAA GATACCAGCA GTAGTCGCCT CTCTTTCAGC AGTTACCCAG 960
 5 GGTTTTGGGA GTCTCTGGAT GATTTTACATA TTCTTAGCAG TGGATTGATA TTGCTGCAGA 1020
 CCACAAACAG TGTGTTTAAT AAAACCCCTGC TAAAGCAGGT AATACCCGAG ACTCTCCTGT 1080
 10 CCTGGCAAAG AGTCCGTGTG GCCAATATGA TGGCAGATAG TGGCAAGAGG TGGGCAGACA 1140
 TCTTTTCAAA ATACAACTCT GGCACCTATA ACAATCAATA CATGGTTCTG GACCTGAAGA 1200
 AAGTAAAGCT GAACCACAGT CTTGACAAAG GCACTCTGTA CATTGTGGAG CAAATTCCTA 1260
 15 CATATGTAGA ATATTCTGAA CAACTGATG TTCTACGGAA AGGATATTGG CCCTCCTACA 1320
 ATGTTCTTTT CCATGAAAAA ATCTACAACT GGAGTGGCTA TCCACTGTTA GTTCAGAAGC 1380
 20 TGGGCTTGGA CTACTCTTAT GATTTAGCTC CACGAGCCAA AATTTTCCGG CGTGACCAAG 1440
 GGAAAGTGAC TGATACGGCA TCCATGAAAT ATATCATGCG ATACAACAAT TATAAGAAGG 1500
 ATCCTTACAG TAGAGGTGAC CCCTGTAATA CCATCTGCTG CCGTGAGGAC CCTGAACTCA 1560
 25 CCTAACCCAA GTCCTTGGAG GTTGTTATGA CACAAAAGGT GGCAGATATY TACCTAGCAT 1620
 CTCAGTACAC ATCCTATGCC ATAAGTGGTC CCACAGTACA AGGTGGCCTC CCTGTTTTTT 1680
 30 GCTGGGACCG TTTCAACAAA ACTCTACATC AGGGCATGCC AGAGGTCTAC AACTTTGATT 1740
 TTATTACCAT GAAACCAATT TTGAACTTG ATATAAAATG AAGGAGGGAG ATGACGGACT 1800
 AGAAGACTGT AAATAAGATA CCAAAGGCAC TATTTTAGCT ATGTTTTTCC CATCAGAATT 1860
 35 ATGCAATAAA ATATATTAAT TTGTCAAAAA AAAAAAAAAA AAA 1903

40 (2) INFORMATION FOR SEQ ID NO: 56:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1869 base pairs
 (B) TYPE: nucleic acid
 45 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:

50 ACAGCTTTTC GGGGCCCCGAG TCGACCCAG CGAAGAGAGC GGGCCCCGGA CAAGCTCGAA 60
 CTCGGGCCGC CTCGCCCTTC CCCGGCTCCG CTCCTCTGTC CCCCTCGGGG TCGCGCGCCC 120
 ACGATGCTGC AGGGCCCTGG CTCGCTGCTG CTGCTCTTCC TCGCCTCGCA CTGCTGCCTG 180
 55 GGCTCGGCGC GCGGGCTCTT CCTCTTTGGC CAGCCCGACT TCTCCTAGAA CCGCANCAAT 240
 TGCAAGCCCA TCCCGGTCAA CCTGCAGCTG TGCCACGGCA TGAATACCA GAACATGCGG 300
 60 CTGCCCAACC TGCTGGGCCA CGAGACCATG AAGGAGGTGC TCGAGCAAGC CGGCGCTTGG 360

ATCCCGCTGG TCATGAAGCA GTGCCACCCG GACACCAAGA AGTTCCTGTG CTCGCTCTTC 420
 5 GCCCCCGTCT GCCTCGATGA CCTAGACGAG ACCATCCAGC CATGCCACTC GCTCTGCGTG 480
 CAGGTGAAGG ACCGCTGCGC CCCGGTCATG TCCGCCTTCG GYTTCCCTTG GCCCGACATG 540
 CTTGAGTGCG ACCGTTTCCC CCAGGACAAC GACCTTTGCA TCCCCCTCGC TAGCAGCGAC 600
 10 CACCTCCTGC CAGCCACCGA GGAAGCTCCA AAGGTATGTG AAGCCTGCAA AAATAAAAAT 660
 GATGATGACA ACGACATAAT GGAAACGCTT TGTAAAAATG ATTTTGCACT GAAAAATAAA 720
 15 GTGAAGGAGA TAACCTACAT CAACCGAGAT ACCAAAATCA TCCTGGAGAC CAAGAGCAAG 780
 ACCATTTACA AGCTGAACGG TGTGTCCGAA AGGGACCTGA AGAAATCGGT GCTGTGGCTC 840
 AAAGACAGCT TGCAGTGAC CTGTGAGGAG ATGAACGACA TCAACGCGCC CTATCTGGTC 900
 20 ATGGGACAGA AACAGGGTGG GGAGCTGGTG ATCACCTCGG TGAAGCGGTG GCAGAAGGGG 960
 CAGAGAGAGT TCAAGCGCAT CTCCCGCAGC ATCCGCAAGC TGCAGTGCTA GTCCCGGCAT 1020
 25 CCTGATGGCT CCGACAGGCC TGCTCCAGAG CACGGCTGAC CATTTCTGCT CCGGGATCTC 1080
 AGCTCCCGTT CCCCAAGCAC ACTCCTAGCT GCTCCAGTCT CAGCCTGGGC AGCTTCCCCC 1140
 TGCCTTTTCG ACGTTTGCAT CCCCAGCATT TCCTGAGTTA TAAGGCCACA GGAGTGGATA 1200
 30 GCTGTTTTCA CCTAAAGGAA AAGCCCACCC GAATCTTGTA GAAATATTCA AACTAATAAA 1260
 ATCATGAATA TTTTATGAA GTTTAAAAAT AGCTCACTTT AAAGCTAGTT TTGAATAGGT 1320
 35 GCAACTGTGA CTTGGGTCTG GTTGGTTGTT GTTTGTTGTT TTGAGTCAGC TGATTTTCAC 1380
 TTCCCACTGA GGTTGTCATA ACATGCAAAT TGCTTCAATT TTCTCTGTGG CCCAAACTTG 1440
 TGGGTCACAA ACCCTGTTGA GATAAAGCTG GCTGTTATCT CAACATCTTC ATCAGCTCCA 1500
 40 GACTGAGACT CAGTGTCTAA GTCTTACAAC AATTCATCAT TTTATACCTT CAATGGGAAC 1560
 TTAAACTGTT ACATGTATCA CATTCCAGCT ACAATACTTC CATTTATTAG AAGCACATTA 1620
 45 ACCATTTCTA TAGCATGATT TCTTCAAGTA AAAGGCAAAA GATATAAATT TTATAATTGA 1680
 CTTGAGTACT TTAAGCCTTG TTTAAAACAT TTCTTACTTA ACTTTTGCAA ATTAAACCCA 1740
 TTGTAGCTTA CCTGTAATAT ACATAGTAGT TTACCTTTAA AAGTTGTAAA AATATTGCTT 1800
 50 TAACCAACAC TGTAATATT TCAGATAAAC ATTATATTCT TGTATATAAA CTTTACATCC 1860
 TGTTTTACC 1869

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(2) INFORMATION FOR SEQ ID NO: 57:

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 1259 base pairs

(B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 57:

	ACCGTGGTTCG TGGGCGGACG GCGGCTGCAG CGYGGAGGAG CTGGGGTTCGC TGTGGGTTCGC	60
	GAACAGAGCC CGGGACGTGC GCGCTTGGTG CACGATCCTG AAGGGGAGCT CCGAGGGGCC	120
10	CGGGTCKCCA GGGCTGCTGC GGCCATTCCC GGAGCCCGGC GCGGGGCCCG NRAGATACTG	180
	GTTTAGGCCG TCCCAGGGCT CCGGGCGCAC CCGKTGGCCG CTGCTGCAGC GGAGGGAGCG	240
15	CGGCGGCGSG NNGGCTCGGA GACAGCGTTT CTCCCGAAT CTCCTCGGG CAGCARGTGG	300
	GAAGTGGGAG CCGGAGCGGC ACTGGCARGC TTCTCTCCGC ANGTCGGCAC CATGCGCCCT	360
	GCAGCCCTGC GCGGGGCCCT GCTGGGCTGC CTCTGCCTGG CGTTGCTTTG CCTGGGCGGT	420
20	GCGGACAAGC GCCTGCGTGA CAACCATGAG TGGAAAAAAC TAATTATGGT TCAGCACTGG	480
	CCTGAGACAG TATGCGAGAA AATTCAAAAC GACTGTAGAG ACCCTCCGGA TTAGTGACAA	540
25	ATACATGGAC TATGGCCCGA TAAAAGTGAA GGATGTAATA GATCGTGGCC CTTCAATTTA	600
	GAAGAGATTA AGGATCTTTT GCCAGAAATG AGGGCATACT GGCCTGACGT AATTCACTCG	660
	TTTCCCAATC GCAGCCGCTT CTGGAAGCAT GAGTGGGAAA AGCATGGGAC CTGCGCCGCC	720
30	CAGGTGGATG CGCTCAACTC CCAGAAGAAG TACTTTGGCA GAAGCCTGGA ACTCTACAGG	780
	GAGCTGGACC TCAACAGTGT GCTTCTAAAA TTGGGGATAA AACCATCCAT CAATTACTAC	840
35	CAAGTTGCAG ATTTTAAAGA TGCCCTTGCC AGAGTATATG GAGTGATACC CAAAATCCAG	900
	TGCCTTCCAC CAAGCCAGGA TGAGGAAGTA CAGACAATTG GTCAGATAGA ACTGTGCCTC	960
	ACTAAGCAAG ACCAGCAGCT GCAAACTGC ACCGAGCCGG GGGAGCAGCC GTCCCCCAAG	1020
40	CAGGAAGTCT GGCTGGCAAA TGGGGCCGCC GAGAGCCGGG GTCTGAGAGT CTGTGAAGAT	1080
	GGCCCACTCT TCTATCCCCC ACCTAAAAAG ACCAAGCATT GATGCCCAAG TTTTGGAAAT	1140
45	ATTCTGTTTT AAAAAGCAAG AGAAATTCAC AAAGTGCAGC TTTCTNAAAA AAAAANAAAA	1200
	AAAAATTGGG GGGTTTTTTT GGGGSGCCCG GGGCCCTTGG TTTTTCCTCC CGGGGGGGT	1259

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(2) INFORMATION FOR SEQ ID NO: 58:

55 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1186 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:

5 CCGCATGGAG AATGGCTCCG CTTCTGTTGC AGCTGGCGGT GCTCGGCGCG GCGCTGGCGG 60
 CCGCAGCCCT CGTACTGATT TCCATCGTTG CATTTACAAC TGCTACAAAA ATGCCAGCAC 120
 TCCATCGACA TGAAGAAGAG AAATTCTTCT TAAATGCCAA AGGCCAGAAA GAAACTTTAC 180
 CCAGCATATG GGA CTCACCT ACCAAACAAC TTTCTGTCGT TGTGCCTTCA TACAATGAAG 240
 10 AAAAACGGTT GCCTGTGATG ATGGATGAAG CTCTGAGCTA TCTAGAGAAG AGACAGAAAC 300
 GAGATCCTGC GTTCACTTAT GAAGTGATAG TAGTTGATGA TGGCAGTAAA GATCAGACCT 360
 15 CAAAGGTAGC TTTTAAATAT TGCCAGAAAT ATGGAAGTGA CAAAGTACGT GTGATAACCC 420
 TGGTGAAGAA TCGTGGAAAA GGTGGAGCGA TTAGAATGGG TATATTCACT TCTCGAGGAG 480
 AAAAGATCCT TATGGCAGAT GCTGATGGAG CCACAAAGTT TCCAGATGTT GAGAAATTAG 540
 20 AAAAGGGGCT AAATGATCTA CAGCCTTGGC CTAATCAAAT GGCTATAGCA TGTGGATCTC 600
 GAGCTCATTT AGAAAAAGAA TCAATTGCTC AGCGTTCTTA CTTCCGTACT CTTCTCATGT 660
 ATGGGTTCCT CTTTCTGGTG TGGTTCCTTT GTGTCAAAGG AATCAGGGAC ACACAGTGTG 720
 25 GGTTCAAATT ATTTACTCGA GAAGCAGCTT CACGGACGTT TTCATCTCTA CACGTTGAAC 780
 GATGGGCATT TGATGTAGAA CTA CTGTACA TAGCACAGTT CTTTAAAATT CCAATAGCAG 840
 30 AAATTGCTGT CAACTGGACA GAAATTGAAG GTTCTAAATT AGTTCCATTC TGGAGCTGGC 900
 TACAAATGGG TAAAGACCTA CTTTTTATAC GACTTCGATA TTTGACTGGT GCCTGGAGGC 960
 TTGAGCAAAC TCGGAAAATG AATTAGGTTG TTTGCAGTCT TCAGTTGTGT TCTTATGCTT 1020
 35 CAGTGTCACT TTTCACTTCA TTTGAACTA AAATTTTAAG TAAAGCTGAA ATAAACTTCT 1080
 TGTCATTGTC TGCCTTTTGA TAATTTTAAA GAAATAACTT TCCATAAGTA AAAAATTATA 1140
 40 TATCTCTTTG GATATAAATG ATTTTAAAA GATGTTTATT TAAAAA 1186

45 (2) INFORMATION FOR SEQ ID NO: 59:

(i) SEQUENCE CHARACTERISTICS:

50 (A) LENGTH: 428 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:

55 GATCCCCCGG CTGCAGGATT CGGCACGAGT ACTGATTCTT TACTAGCTTT TTTTAGTATA 60
 AGCAGAGTTC CAAGTCTCCC CTAGGGTTGT CTCTACATTT TTTTACATTT CCAGTGGGTA 120
 60 RGGTTTAGCT GGGGAAGGA CATTTTATAA GGGTTAGTTT TTTTATTTAG TATGGACATT 180

TGCTTTTTC ATTACGTACT GTTGTTTTC CTTGTTAGGT GTGCTTTGGT GGTTTTAATA 240
 TTATTGTGCC AGGGATGGGG AAATGGGGG GGTGTGTGG GAAGAGTACT TATTATTGTG 300
 5 ...TCTTCAG TGTAAATGTT CTTGGTAATT GATACCTCTC TGTTTTATTT NTCTCATCT 360
 TCAAAATAA AACTTTTTGA AATTTGAAAA AAAAAAAAAA NAAAAAATC GGGGGGGGGC 420
 CCGGTACC 428
 10

15 (2) INFORMATION FOR SEQ ID NO: 60:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 501 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 20 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:

GGCACGAGCT TTCAGCAGGG GACAGCCCGA TTGGGGACAA TGGCGTCTCT TGGCCACATC 60
 25 TTGTTTTCT GTGTGGGTCT CCTCACCATG GCCAAGGCAG AAAGTCCAAA GGAACACGAC 120
 CCGTTCACCTT ACGACTACCA GTCCCTGCAG ATCGGAGGCC TCGTCATCGC CGGGATCCTC 180
 30 TTCATCCTGG GCATCCTCAT CGTGCTGAGC AGAAGATGCC GGTGCAAGTT CAACCAGCAG 240
 CAGAGGACTG GGAACCCGA TGAAGAGGAG GGAAGTTTCC GCAGCTCCAT CCGCCGCTCTG 300
 TCCACCCGCA GCGGGTAGAA ACACCTGGAG CGATGGAATC CCGCCAGGAC TCCCCTGGCA 360
 35 CCTGACATCT CCCACGCTCC AACTGCGCGC CCACCGCCCC CTCCGCGGCC CCTTCCCCAG 420
 CCCTGCCCCC GCAGACTCCC CCTGCCGCCA AGACTTCCAA TAAAACGTGC GTTCCTCTCG 480
 40 AAAAAAAAAA AAATAAAAAA A 501

45 (2) INFORMATION FOR SEQ ID NO: 61:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1197 base pairs
 (B) TYPE: nucleic acid
 50 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:

ACATGATGGN TACCAAAGAA TTCGGCANAG GGCGCGCACT TCAATATCG 60
 AGTGCCTGCG GGAAGTCTG ACGCCCCCGC TGCTGTCTCT TACGTGGGCG 120
 60 CCCCCAGGC CCTCACCTG AAGCTCCCAG TGACCAKCAA CAGCCCACCG 180

207

AGATGGCGGC CCAGGATTTC TTCCAGCGCT GGAAGCAGCT GAGCCTCCCT CAACAGGAGG 240
 CGCAGAAAAT CTTCAAAGCC AACCACCCCA TGGACGCAGA AGTTACTAAG GCCAAGCTTC 300
 5 TGGGGTTTGG CTCTGCTCTC CTGGACAATG TGGACCCCAA CCCTGAGAAC TTCGTGGGGG 360
 CGGGGATCAT CCAGACTAAA GCCCTGCAGG TGGGCTGTCT GCTTCGGCTG GAGCCCAATG 420
 10 CCCAGGCCCA GATGTACCGG CTGACCCTGC GCACCAGCAA GGAGCCCGTC TCCCCTCACC 480
 TGTGTGAGCT GCTGGCACAG CAGTTCTGAG CCCTGGACTC TGCCCCGGGG GATGTGGCCG 540
 GCACTGGGCA GCCCCCTGGA CTGAGGCAGT TTTGGTGGAT GGGGGACCTC CACTGGTGAC 600
 15 AGAGAAGACA CCAGGGTTTG GGGGATGCCT GGGACTTTCC TCCGGCCTTT TGTATTTTTA 660
 TTTTGTTC A TCTGCTGCTG TTTACATTCT GGGGGGTTAG GGGGAGTCCC CCTCCCTCCC 720
 TTTCCCCCCC AAGCACAGAG GGGAGAGGGG CCAGGGAAGT GGATGTCTCC TCCCCTCCCA 780
 20 CCCCACCTG TTGTAGCCCC TCCTACCCCC TCCCCATCCA GGGGCTGTGT ATTATTGTGA 840
 GCGAATAAAC AGAGAGACGC TAACAGCCCC ATGTCTGTGT CCATCACCCA CTGTTAGGTA 900
 25 GTCAAAGAAG TGGGGTGAGG GCATGCAGAG TGTGGGTGGC CAGNTTCGCA GCCCATGGGT 960
 GGGACTCTGG GGAGACAGCA GCAGCAGCAG CCGCCGAAGC CCCAGCTGCA AGGCCACCAG 1020
 ACGCACTCCT GTGCCTGGTT CTTYAGTCCC CAACACCAGG TAGCAAGCTY TGGGCAGCTG 1080
 30 GGCCTGGTAG ACCTCATCTT CTGTCTTCTY TGGTGGCCCT GGCTCTGGTG GGAAGTGCGT 1140
 GGAGGTGACC AGGGTATAGA AGTTTCGGAG CTGATTGGAA GAGGATTAAC TTCCCGC 1197
 35

(2) INFORMATION FOR SEQ ID NO: 62:

40 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 595 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:

ATTNANGACK TKYAGCCTYT WATACMATCA TTATAGGGAR AAGCTGGTAC GCCTGMARGT 60
 50 ACCGGTCYGG AATTCNCGGG TCGACCCACG CGTCCGGCAC AGCGGGAGTT GGTCTTGACA 120
 CCAGATGTTT TCTGCTCCTG GTTAATGTCA GTGAGGGCTG GAAGTTGAAT AAATGAGAAC 180
 AGGAGTGGTC TGGGCCCCATG TAAATGATCC TCCCTTGAAA GGAGGAACAG CTTCATCAT 240
 55 TTGTTCCAGC TAAGCCTTGC ATGCATTATA GATCTGGTGC TAAGCAGTGG GAAAGATCTC 300
 ATAAGTAATG TTTTATGTTT TTTCTGTCTC TCCTCTTCTG TTGTTCTTCG CTTGTGGGTT 360
 60 GTGTTTGTGT GTTAACTGGA AAATTGCTAT AAGCCAGTTG TCTCTAAGTT TTAAAAACGA 420

	ATTAGAAAAA CCATAAAATC TCTGGCCTAT GCACATTGTC CCTGTTTGTG TAAAAACATTA	480
5	AAGGGTAAAT AAAAAGGAAG GAGAACAGTC AATAATGTGC ATCAAATATA TTCTGAGTTC	540
	TAGAGAAATT AATGACCAAG CATTAGAACT AGAAGCAAAA AAAAAAAAAA AAAAA	595
10	(2) INFORMATION FOR SEQ ID NO: 63:	
	(i) SEQUENCE CHARACTERISTICS:	
15	(A) LENGTH: 1478 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63:	
	CGGCGCTGAG GACGCACGGA TGCCTTCCGT GCCTTCCATC AAGATCTCAA TTTGTGCGC	60
	AAGTTCCTAC AGCCCCCTGTT GATTGGAGAG CTGGCTCCGG AAGAACCCAG CCAGGATGGA	120
25	CCCCGAATG CGCATGGTCG AGGACTTCCG AGCCCTGCAC CAGGCAGCCG AGGACATGAA	180
	GCTGTTTGAT GCCAGTCCCA CCTTCTTTGC TTTCTACTG GGCCACATCC TGGCCATGGA	240
	GGTGTGCGCC TGGCTCCTTA TCTACCTCCT GGTCTCTGGC TGGGTGCCCC GTGCCCTGGN	300
30	CCGCCTTCAT CCTGGCCATC TCTCAGGCTC AGTCCTGGTG TCTGCAGCAT GACCTGGGCC	360
	ATGCTCCATC TTCAAGAAGW CCTGGTGGAA CCACGTGGCC CAGAAGTTCG TGATGGGGCA	420
35	GCTAAAGGGC TTCTCCGCCC ACTGGTGGAA CTTCGCCCAC TTCCAGCACC ACGCCAAGCC	480
	CAACATCTTC CACAAAGACC CAGACGTGAC GGTGGCGCCC GTCTTCTCTC TGGGGGAGTC	540
40	ATCCGTCGAG TATGGCAAGA AGAAACGCAG ATACCTACCC TACAACCAGC AGCACCTGTA	600
	CTTCTTCCTG ATCGGCCCCG CGCTGCTCAC CCTGGTGAAC TTTGAAGTGG AAAATCTGGC	660
	GTACATGCTG GTGTGCATGC AGTGGGCGGA TTTGCTCTGG GCCGCCAGCT TCTATGCCCC	720
45	CTTCTTCTTA TCCTACCTCC CCTTCTACGG CGTCCCTGGG GTGCTGCTCT TCTTTGTGTC	780
	TGTCAGGGTC CTGGAAAGCC ACTGGTTCGT GTGGATCACA CAGATGAACC ACATCCCCAA	840
50	GGAGATCGGC CACGAGAAGC ACCGGGACTG GGTGAGCTCT CAGCTGGCAG CCACCTGCAA	900
	CGTGGAGCCC TCACTTTTCA CCAACTGGTT CAGCGGGCAC CTCAACTTCC AGATCGAGCA	960
	CCACCTCTTC CCCAGGATGC CGAGACACAA CTACAGCCGG GTGGCCCCGC TGGTCAAGTC	1020
55	GCTGTGTGCC AAGCACGGCC TCAGCTACGA ATGAAGCCCT TCTCACCAGC GCTGGTGGAC	1080
	ATCGTCAGGT CCCTGAAGAA GTCTGGTGAC ATCTGGCTCG AGGCCTACCT CCATCAGTGA	1140
60	AGGCAACACC CAGGCGGGCA GAGAAGGGCT CAGGGCACCA GCAACCAAGC CAGCCCCCGG	1200

CGGGATCGAT ACCCCCACCC CTCCACTGGC CAGCCTGGGG GTGCCCTGCC TGCCCTCCTG 1260
 GTACTGTTGT CTTCCCCTCG GCCCCCTCAC ATGTGTATTC AGCAGCCCTA TGGCCTTGGC 1320
 5 TCTGGGCCTG ATGGGACAGG GGTAGAGGGA AGGTGAGCAT AGCACATTTT CCTAGAGCGA 1380
 GAATTGGGGG AAAGCTGTTA TTTTATATATT AAAATACATT CAGATGTAAA AAAAAAAAAA 1440
 AAAAACTCGA GGGGGGGCCC CGGNAACCAA TTCGCCCT 1478
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(2) INFORMATION FOR SEQ ID NO: 64:

15

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2033 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

20

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 64:

GGCAGGAGGA AGAACGCAAA GCTGAGAACA TGGACGTTAA TATCGCCCCA CTCCGCGCCT 60
 25 GGGACGATTT CTTCCCGGGT TCCGATCGCT TTGCCCGGCC GGACTTCAGG GACATTTCCA 120
 AATGGAACAA CCGCGTAGTG AGCAACCTGC TCTATTACCA GACCAACTAC CTGGTGGTGG 180
 30 CTGCCATGAT GATTTCCATT GTGGGGTTTC TGAGTCCCTT CAACATGATC CTGGGAGGAA 240
 TCGTGGTGGT GCTGGTGTTC ACAGGGTTTG TGTGGGCAGC CCACAATAAA GACGTCCTTC 300
 GCCGGATGAA GAAGCGCTAC CCCACGACGT TCGTTATGGT GGTTCATGTTG GCGAGCTATT 360
 35 TCCTTATCTC CATGTTTGGG GGAGTCATGG TCTTTGTGTT TGGCATTACT TTTCCTTTGC 420
 TGTGATGTT TATCCATGCA TCGTTGAGAC TTCGGAACCT CAAGAACAAA CTGGAGAATA 480
 40 AAATGGAAGG AATAGGTTTG AAGAGGACAC CGATGGGCAT TGTCTGGAT GCCCTAGAAC 540
 AGCAGGAAGA AGGCATCAAC AGACTCACTG ACTATATCAG CAAAGTGAAG GAATAACAT 600
 AACTTACCTG AGCTAGGGTT GCAGCAGAAA TTGAGTTGCA GCTTGCCCTT GTCCAGACCT 660
 45 ATGTTCTGCT TGCCTTTTGG AAACAGGAGG TGCACGTACC ACCCAATTAT CTATGGCAGC 720
 ATGCATGTAT AGGCCGAAC ATTATCAGCT CTGATGTTTC AGAGAGAAGA CCTCAGAAAC 780
 50 CGAAAGAAAA CCACCACCCT CCTATTGTGT CTGAAGTTTC ACGTGTGTTT ATGAAATCTA 840
 ATGGGAAATG GATCACACGA TTTCTTTAAG GGAATTAAAA AAAATAAAAG AATTACGGCT 900
 TTTACAGCAA CAATACGATT ATCTTATAGG AAAAAAAAAAT CATTTGTAAG TATCAAGACA 960
 55 ATACGAGTAA ATGAAAAGGC TGTAAAGTA GATGACATCA TGTGTTAGCC TGTTCCTAAT 1020
 CCCCTAGAAT TGTAATGTGT GGGATATAAA TTAGTTTTTA TTATCTCTTT AAAAAATCAA 1080
 60 GATGATCTCT ATCACTTTGC CACCTGTTTG ATGTGCAGTG GAAACTGGTT AAGCCAGTTG 1140

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TTCATACTTC CTTTAC... TAAAGATAG CTGTTTAGGA TATMTTGTTA CATTTTGTGA 1200
 AATTTTGTGA ATGCTAGTAA TGTGTTTCA CCAGCAAGTA TTTGTTGCAA ACTTAATGTC 1260
 ATTTTCCTTA AGATGGTTAC AGCTATGTAA CCTGTATTAT TCTGGACGGA CTTATTAAAA 1320
 TACAAACAGA CAAAAAATAA AACAAACTT GAGTTCTATT TACCTTGCAC ATTTTGTGTT 1380
 GTTACAGTGA AAAAAATGGT CCAAGAAAAT GTTGCCATT TTTGCATTGT TTCGTTTMTA 1440
 ACTGGAACAT TTAGAAAGAA GGAAATGAAT GTGCATTTTA TTAATTCCTT AGGGGCACAA 1500
 GGAGGACAAT AATAGCTGAT CTTTGAAAT TTGAAAAACG TCTTTAGATG ACCAAGCAAA 1560
 AAGCTTTAAA AAATGGTAAT GAAAATGGAA TGCAGCTACT GCAGCTAATA AAAAATTTTA 1620
 GATAGCAATT GTTACAACCA TATGCCTTTA TAGCTAGACA TTAGAATTAT GATAGCATGA 1680
 GTTTATACAT TCTATTATTT TTCTCCCTT TCTCATGTTT TTATAAATAG GTAATAAAAA 1740
 ATGTTTGGCC TGCCAATTGA ATGATTTTCGT AGCTGAAGTA GAAACATTTA GGTTCCTGTA 1800
 GCATTAAATT GTGAAGACAA CTGGAGTGGT ACTTACTGAA GAACTCTCT GTATGTCCTA 1860
 GAATAAGAAG CAATGATGTG CTGCTTCTGA TTTTCTTGC ATTTTAAATT CTCAGCCAAC 1920
 CTACAGCCAT GATCTTTAGC ACAGTGATAT CACCATGACT TCACAGACAT GGTCTAGAAT 1980
 CTGTACCCTT ACCCACATAT GAAGAATAAA ATTGATTAAA GGTAAAAAA AAA 2033

35 (2) INFORMATION FOR SEQ ID NO: 65:

(i) SEQUENCE CHARACTERISTICS:

40 (A) LENGTH: 440 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:

45 ATGTTTCTTA CTAGAATACT GTGTCCAACC TATATAGCCC TAACTTTCCT GGTTCACATT 60
 GTGGCCCTAG TATCTGGGCA GCTGTGCATG GAGATAGCCA GAGGAAACAT TTTTCTCTT 120
 AATGAATTGG TGACCACATT TTGTTGTTCT TGCCTCCTAT TATCCGTGCC CTATTTGCAT 180
 50 CCTGGTTTCT TCTACAGTAG TTTATGTAAA TGTGTTTTG TCCTTGTCGT TCTCAGTAGA 240
 ATTGGTTCTG TAAACGAAAC CTGGTCCTGT AATTTTCAGTA TATGCTCATA TCTCATCTTT 300
 55 GGCTCTCCCA TTTCACAGC AGTGATCCCT AAAAGATGTG CCCTAGAGGA TATCCAGAAC 360
 AATCCAATTG GATGTCTTCT CCGCTGCACT CCAGCCTGGG AGACAGAGGG AGACTCNATC 420
 60 TCAAAAAAAA TAAAAAAA 440

(2) INFORMATION FOR SEQ ID NO: 66:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 3301 base pairs

(B) TYPE: nucleic acid

10

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:

15	GGTCATAAGG GGAGGGTTGN NGTGTGTCCC TCCAGGTTGT GCAGAGGGGA TTAGAAGTAA	60
	GTAGGTTAGA GGGGAGGTGG AGGGAGTGTG CTGGGTGTG AGCTTTTATG ATGCTGAAAG	120
	GATCATGATA TGCTAAGGAC AGGATAGTGT TGGGTGTGAC ACACAGGTGT AGGCAATCCT	180
20	GGTGGCTAGT ATGTAAAAGT GAATGTCCTG ACTCCCTTAG AGGGTACCTG NCAGAGTGCC	240
	CTTGARGGA CTAGTGCTGG AGAAATTAAT AGGAGAGGGG ACGGGCATCC ATTAACCTTT	300
	TCTTGCTGC AGCCTGTAGG GTCCAGCGTC AAAGCGAATC ATGGGGTCCA GGGCTGAGCT	360
25	GTGCACTCTC TTAGGCGGAT TCTCCTTCCT CCTGCTACTG ATACCAGGCG AGGGGGCCAA	420
	GGGTGGATCC CTCAGAGAGA GTCAGGGAGT CTGCTCCAAG CAGACACTGG TGGTCCCCT	480
30	CCACTACAAC GAGTCCTACA GCCAACCAGT GTACAAGCCC TACCTGACCT TGTGCGCTGG	540
	GAGCGCATCT GCAGCACTTA CAGGACCATG TACCGCGTTA TGTGGCGGGA GGTGAGGCGG	600
	GAGGTTGAGC AGACCCATGC AGTGTGCTGC CAGGGCTGGA AGAAGCGGCA CCCGGGGGCG	660
35	CTCACCTGTG AAGCCATCTG CGCCAAGCCT TGCCTGAACG GAGGCCTCTG CGTTAGGCCT	720
	GACCAGTGCG AGTGCGCCCC CGGCTGGGGA GGAAGCACT GTCATGTGGA CGTGGATGAA	780
40	TGTAGGACCA GCATCACCTT CTGCTCGCAC CATTGTTTTA ATACGGCARG CAGCTTCAMC	840
	TGCGGCTGCC CCATGACCTA GTGCTAGGCG TGGACGGGCG CACCTGCATG GAGGGGTCCC	900
	CAGAGCCCCC AACCAGTGCC AGCATACTCA GCGTGGCCST TCGGGARGCG GAAAAAGATG	960
45	ACGCGCTCTG AAGCAGGAGA TTCACGAGCT GCGAGGCCCT TGAAGCGGCT GGAGCAGTGG	1020
	NCCGGTCAGC TGGGCCCTGG NTCAGACGGT GCTGCCCCGT CCGCCTGAAG WGCTGCAGCC	1080
50	AGAACAGGTG GCTGAGCTGT GGGGCCGGGG TGACCGGATC GAATCTCTCA GCGACCAGGT	1140
	GCTGCTGCTG GAGGAGAGGC TAGGTGCCTG CTCCTGTGAG GACAACAGCC TGGGCCTCGG	1200
	CGTCAATCAT CGATAAGAAG CCTCTACAGC ACCCCTGCCC CCTAATTTAT ACAGAAACCG	1260
55	GACCCACTAA TCCTCTGGGA TTGGCCGACT GTGAGCTGCA GATAAGGCTA TCAGCCACCA	1320
	AAGAGCAATG AACAATGGAA ACTTCAGAGA GCTGAAGAAA GCGGTAGGCG TGTGTTCTTG	1380
60	GCCTGCCCTT GAGTCTTCTG GCTGGGGGCA GGTGCTCTGG GCAAGAACTG CTTCTTCAAT	1440

	TCCTTAACAA ATGCAACCAC CAACACCCAG ATCTCTCTCT CTCTTTATTT TCAGTTTTTT	1500
5	TGCTGTTATC CAGATAATTA ATAAAAACCA ACCACGCAA ACTGGGTCCC ACCCTCTCCT	1560
	TTT CTCCCA GCCTACCTCC CCAGTTGTGG GAACAGGTCT GGAGTGAGAG GCAGGGAGTG	1620
	GCTAATGCCN CCAGGAAGAA ATGAAAAC TGCTCAGAGAG GGGGAAGCCT CAACAGAAAA	1680
10	AGAAATAAAT TAAAAGCCCT CCTATCCCTT CCAGCCAGGG TTCGTTCCCT TCCCCAACTC	1740
	CCCAGGGGGC AGAAGTGAGT GCAGCACCTG ATGTCTGCTT CTTCCTTGTG TGTCTGGTGA	1800
15	GATGGTGCAG CAGGGCTGCA GGGGGCTGGG TGGGGTCATG TCCACTGAAG AACTGTACTA	1860
	TGGGGACAGA AAACCAGAAA TGTGGAGACT GAACTGGTAT CCCAGAGAGT GCACGACCTT	1920
	GGGCATCTGG GCAAGGGCAG GCATGAGACC TCTGAATTAG AAGGGTCCAG CCCCCACTGA	1980
20	CAGGAGGCTA CACTGGGAGG GAAGGTGAAG GTGCTGAGGA AAGCTCCCAT GATGAGCCTG	2040
	GGAGTGCTTC AGGTATCAGC TTCCAGCCAG AGGGCGAGAA GTCTCTCTCA CAAATGGATG	2100
25	AGTCCATTGA ATCCATGGAC TTTGGAGTGG GGGGGATTG TTCCAAAGAA TGGATGAGTC	2160
	CACTGGCCAA TGTGGGGTAG AGGGGTAGAG AAGACCACAT AGGAAGAGAC TCCACTGGGG	2220
	ATGGAATGTT CCCCTCCCTT GTGTAGGCTG AGTCACTGGA GATGAGGGGG AGGCAACTGT	2280
30	CCCACAGACA ARACAGTAGG AGGTGGGGGT CAAGAGTGGA GACTGCACCG AGGCAAGAGT	2340
	CCATGGATGG GGCCAAGAGG GGGCAGGAGT GGCGCTGTAT CCACATTTCA CTTCAGAAGT	2400
35	TGAAGATTCC AAAGAGGAGA ATAAGTGGGG AGAGGGGAGA CAAGGAAGAG GGT'TTKGCCC	2460
	TGCTTCAGGG CCCACTGGGT GGGTAGGTGT GGGGAGGAAG ATGGGGACAG ATGGGAGGAG	2520
	AGCTCAGAGC CAGGGTTCAC CCACCGCCCC CAGGCTTCTT CAGATAGTCA CCACCACCCC	2580
40	GGCCATCAGT GGAGATTTCC CGGAAAACAG TGAAGCATGG AGTGCCGGAC TCTGTGAGCC	2640
	AGAGCTGGGA CGTCATCTGG TGTGAGCCCT TCCGTGGGCA CTGGGGGCAG CACCCGACCC	2700
45	TGACATTGTC CCGAGGTGAA GCGACGCTCC TTCTTGCACT AGAAGTCTTG GTAGGAGGAC	2760
	ATGACTATGG GGACAATGGG AACCTGGGCC TGCCTGCAA GATGGAAGGC GCCACGTTTG	2820
	AAGGGCAGCA TGGAGCCATT GTGGTTTCTC GTTCCCTCAG GAAACACCCA GACCYTCACG	2880
50	TCCTGGGTGA GCAGGTCTG GGCGACCTCA GACATGACAC TGATGGCATC CCCCCTGCGC	2940
	TTCCGGTCTGA TGAAGATGAC TCCTGCCAGC CAGCAGGCCA GCGCGCAGAG CCAGCCCACA	3000
55	GTANTCGCGC TTGGCAATGG GCACACAGCG GCCTGGCAGT ACCTCCATCA TCCCAAGCAG	3060
	ATCGAGAGAG CTCTGGTGGT TGGAGACAAC AACATAGGGC TGGGAGGAG GGAAGTGGTG	3120
	AGCCCCTCGC ACCTCCACTC GGATCCCGTA CAGGTATTTG ATGTGGAGCA GCATTAGACG	3180
60	CAAGATCTTC ATGTTCTCGA CGTTGCGTCC TCGCACGGCA CACACAGAGA TGGCGAGCAC	3240

AGCCAGGAAG AGGATCCAGC CATTGTAGAA GGCCATCTTG AAGAAGTACT TGGCACTGGG 3300

G 3301

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(2) INFORMATION FOR SEQ ID NO: 67:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1535 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

15

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:

20 GGCACGAGGT CAAGCGAAAG GATTTCAAGG AACAGATCAT CCACCATGTG TTCACCATCA 60
 TTCTCATCAG CTTTTCCTGG TTTGCCAATT ACATCCGAGC TGGGACTCTA ATCATGGCTC 120
 TGCATGGACT CTTCGGATTA CCTGCTGGAG TCAGCCAAGA TGTTTAACTA CGCGGGATGG 180
 25 AAGAACACCT GCAACAACAT CTTTCATCGTC TTCGCCATTG TTTTATCAT CACCCGACTG 240
 GTCATCCTGC CCTTCTGGAT CCTGCATTGC ACCCTGGTGT ACCCACTGGA GCTCTATCCT 300
 GCCTTCTTTG GCTATTACTT CTTCAATTCC ATGATGGGAG TTCTACAGCT GCTGCATATC 360
 30 TTCTGGGCCCT ACCTCATTTT GCGCATGGCC CACAAGTTCA TAACTGGAAA GCTGGTAGAA 420
 GATGAACGCA GTACCGGGAA GAAACAGAGA GCTCAGAGGG GGAGGAGGCT GCAGCTGGGG 480
 35 GAGGAGCAAA GAGCCGGCCC CTAGCCAATG GCCACCCCAT CCTCAATAAC AACCATCGTA 540
 AGAATGACTG AACCATTATT CCAGCTGCCT CCCAGATTAA TGCATAAAGC CAAGGAACTA 600
 CCCCCTCCC TGCGCTATAG GGTCACTTTA AGCTCTGGGG AAAAAGGAGA AAGTGAGAGG 660
 40 AGAGTTCTCT GCATCCTCCC TCCTTGCTTG TCACCCAGTT GCCTTTAAAC CAAATTCTAA 720
 CCAGCCTATC CCCAGTAGG GGGACGTTGG TTATATTTCTG TTAGAGGGGG ACGGTCGTAT 780
 45 TTTCTCCCT ACCCGCAAG TCATCCTTTC TACTGCTTTT GAGGCCCTCC CTCAGCTCTC 840
 TGTGGGTAGG GGTTACAATT CACATTCCTT ATTCTGAGAA TTTGGCCCCA GCTGTTTGCC 900
 TTTGACTCCC TGACCTCCAG AGCCAGGGTT GTGCCTTATT GTCCCATCTG TGGGCCTCAT 960
 50 TCTGCCAAAG CTGGACCAAG GCTAACCTTT CTAAGCTCCC TAACTTGGGC CAGAAACCAA 1020
 AGCTGAGCTT TTAACCTTCT CCCTCTATGA CACAAATGAA TTGAGGCTAG GACGAGGGTG 1080
 55 CACATAACCC TTACCCTACC TCTGCCAAAA AGTGGGGGCT GGAATGAGGA CTGCTCGGAT 1140
 GATCTTTCTT AGTGCTACTT CTTTCAGCTG TCCCTGTAGC TACAGCTCTA AGATCTGACT 1200
 GCCTCCTCCT TTCTCTGGCC TCTTCCCCCT TCCCTCTTCT CTTTACGCTAG GCTAGCTGGT 1260
 60

TTGGAGTAGA ATGGCAACTA ATTCTAATTT TTATTTATTA AATATTTGGG GTTTTGGTTT 1320
 TAAAGCCAGA ATTACGGCTA GCACCTAGCA TTTCAGCAGA GGGACCATTT TAGACCAAAA 1380
 5 TGTACTGTTA ATGGGTPTTTT TTTTAAAATT AAAAGATTAA ATAAAAATA TTAAATAAAA 1440
 CATGCAATA AGTGTGAGAC TATTAGGAAT TGAGAAGGGG GATCAACTAA ATAAACGAAG 1500
 10 AGAGTCTTTC TTATGCAAAA AAAAAAAAAA AAAAA 1535

(2) INFORMATION FOR SEQ ID NO: 68:

15

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1244 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

20

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:

25 GGGCACCCAC CAGCGGCGCC GACCTCAGCG CGCACCTATG GGCTCGCTAC CAGGACATGC 60
 GGAGACTGGT GCACGACCTC CTGCCCCCG AGGTCTGCAG TCTCCTGAAC CCAGCAGCCA 120
 TCTACGCCAA CAACGAGATC AGCCTGCGTG ACGTTGAGGT CTACGGCTTT GACTACGACT 180
 30 ACACCCTGGC CCAGTATGCA GACGCACTGC ACCCCGAGAT CTTCAGTACC GCCCGTGACA 240
 TCCTGATCGA GCACTACAAG TACCCAGAAG GGATTCGGAA GTATGACTAC AACCCCAGCT 300
 35 TTGCCATCCG TGGCCTCCAC TATGACATTC AGAAGAGCCT TCTGATGAAG ATTGACGCCT 360
 TCCACTACGT GCAGCTGGGG ACAGCCTACA GGGGCCTCCA GCCTGTGCCA GACGAGGAGG 420
 TGATTGAGCT GTATGGGGGT ACCCAGCACA TCCCACTATA CCAGATGAGT GGCTTCTATG 480
 40 GCAAGGTCC CTCCATTAAG CAGTTCATGG ACATCTTCTC GCTACCGAG ATGGCTCTGC 540
 TGTCTGTGT GGTGGACTAC TTTCTGGGCC ACAGCCTGGA GTTTGACCAA GCACATCTCT 600
 ACAAGGACGT GACGGACGCC ATCCGAGACG TGCATGTGAA GGGCCTCATG TACCAGTGGA 660
 45 TCGAGCAGGA CATGGAGAAG TACATCCTGA GAGGGGATGA GACGTTTGCT GTCCTGAGCC 720
 GCCTGGTGGC CCATGGGAAA CAGCTGTTCC TCATCACCAA CAGTCCTTTC AGCTTCGTAG 780
 50 ACAAGGGGAT GCGGCACATG GTGGGTCCCG ATTGGCGCCA CTCTTCGATG TGGTCATTGT 840
 CCAGGCAGAC AAGCCCAGCT TCTTCACTGA CCGGCGCAAG CTTTNCAGAA AACTCGATGA 900
 GAAGGGCTCA CTTCAGTGGG ACCGGATCAC CCGCTTGGAA AAGGCGAAGA TATATCGGCA 960
 55 GGGAAACCTG TTTGACTTCT TACGCTTGAC GGAATGGCGT GGGCCCCGCT TGCTCTACTT 1020
 CGGGGACCAC CTCTATAGTG ATCTGGCGGA TCTCATGCTG CCGGACCGCT CCGGCACAGG 1080
 60 CGCCATCATC CCCGAGCTGG AGCGTGAGAT CCGCATCATC AACACGGAGC AGTACATGCA 1140

	CTCGCTKACG TGGCAGCAGG CGCTCACGGG GCTKCTKGAG CGCATKCAGA CCTATCAGGA	1200
5	CGCGGAGTTG AGGCAGGTCT TGCTTCCTTG ATGAAAGANC GNNT	1244
(2) INFORMATION FOR SEQ ID NO: 69:		
10	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1292 base pairs	
	(B) TYPE: nucleic acid	
15	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 69:		
20	GGCACGAGCA GCGACGCGAC TCTGGTGCGG GCCGTCTTCT TCCCCCGAG CTGGGCGTGC	60
	GCGGCCGCAA TGAAGTGGGA GCTGCTGCTG TGGCTGCTGG TGCTGTGCGC GCTGCTCCTG	120
	CTCTTGGTGC AGCTGCTGCG CTTCTTGAGG GCTGACGGCG ACCTGACGCT ACTATGGGCC	180
25	GAGTGGCAGG GACGACGCCC AGAATGGGAG CTGACTGATA TGGTGGTGTG GGTGACTGGA	240
	GCCTCGAGTG GAATTGGTGA GGAGCTGGCT TACCAGTTGT CTAAACTAGG AGTTTCTCTT	300
30	GTGCTGTCTAG CCAGAAGAGT GCATGAGCTG GAAAGGGTGA AAAGAAGATG CCTAGAGAAT	360
	GGCAATTAA AAGAAAAAGA TATACTTGTT TTGCCCTTG ACCTGACCGA CACTGGTTCC	420
	CATGAAGCGG CTACCAAAGC TGTTCTCCAG GAGTTTGGTA GAATCGACAT TCTGGTCAAC	480
35	AATGGTGGAA TGTCCCAGCG TTCTCTGTGC ATGGATACCA GCTTGGATGT CTACAGAAAG	540
	CTAATAGAGC TTAAGTACTT AGGGACGGTG TCCTTGACAA AATGTGTTCT GCCTCACATG	600
40	ATCGAGAGGA AGCAAGGAAA GATTGTTACT GTGAATAGCA TCCTGGGTAT CATATCTGTA	660
	CCTCTTTCCA TTGGATACTG TGCTAGCAAG CATGCTCTCC GGGGTTTTTT TAATGGCCTT	720
	CGAACAGAAC TTGCCACATA CCCAGGTATA ATAGTTTCTA ACATTTGCCC AGGACCTGTG	780
45	CAATCAAATA TTGTGGAGAA TTCCCTAGCT GGAGAAGTCA CAAAGACTAT AGGCAATAAT	840
	GGAGACCAGT CCCACAAGAT GACAACCAGT CGTTGTGTGC GCTTGATGTT AATCAGCATG	900
50	GCCAATGATT TGAAAGAAGT TTGGATCTCA GAACAACCTT TCTTGTCTTAG TAACATATTT	960
	GTGGCAATAC ATGCCAACCT GGGCCTGGTG GATAACCAAC AAGATGAGCA AGAAAAGGAT	1020
	TGAGAAGTTT AAGAGTGGTG TGATGCAGA CTCTCTTATC TCTAAGATCT TTAAGACAAA	1080
55	ACATGACTGA AAAGAGCACC TGTACTTTTC AAGCCACTGA AAGAGGATC TTAAGACATG	1140
	AAAACAGCAA TCTTCTTATG CTTCTGAATA ATCAAAGACT AATCTGATC TTTACTTTTTT	1200
60	AATAGATATG ACTTTGCTTC CAACATGGAA TGAAATAAAA AATAAATAT AAAAGATTGC	1260

CATGAATCTT GCAAAAAAAAA AAAAAAAAAA AA

1292

5

(2) INFORMATION FOR SEQ ID NO: 70:

(i) SEQUENCE CHARACTERISTICS:

10

- (A) LENGTH: 1031 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70:

15

GGGCTGTTGC TTTTGAACAG AACCTATAT TACTCTCCTG GGATCTGAGT TTCTGCAGGT 60

CATTTGTATG TAGGACCAGG AGTATCTCCT CAGGTGACCA GTTTTGGGGA CCCGTATGTG 120

20

GCAAATTCTA AGCTGCCATA TTGAACATCA TCCCCTGGG AGTGGTTATG TTGTATCCCC 180

ATCTTGGCTG GCTTCAGTTT TTGCTGTAGC CCTAGAGCAC TTGTTTGTG GGAGGCTGGC 240

25

CTCTTGCCTA CCTCCTTGCA TGGACAGGGG GATGAATATT TACTTTCCCA CCTCCTTGCT 300

TTTCTTTTCA CTGATACCAC TGAATGGAAC TGGTGCTGTG ACTCCTGCTG CTGGGGATTT 360

ATGTCCCGAG ACCTTAGCCT GGCTGAGTGG AGCCTGAGAC CTGCACAACA GCTCATGGTC 420

30

ATGCATGARA GAGAAGTGGC TGGCCACAGC AGAGGGAACA GTAACAGCCC AGGGGCCTTT 480

ATTTTGGGAA AGGCTGTCCG GGGCTGTTAC TGTCTCTTCT GGTATAAAG CAGACATGTG 540

35

GCCATCTTTT CCGCAGGTTA GAGTGGGCTC CTTTCTTTTT GGAATCCTTT TCTTCTCCTT 600

TGGTAGCAGC TCCCTGCCTC CAGGGCTTCC GCCACCAGCG TCTCTGCTGT GTTGCGCAGT 660

GCAGTGGGGT GCAAGGGCTT TGTTCCTGCC TGCCTGAAAG AGAGGGCTCT GGGGATGGAG 720

40

ATGAGAAACA ACACGCTCTC CTTCAGACAA TGAGGCATTC TGCTCTCCTG CTGCCATTCT 780

TCATCTCCAC TGAGAGCCAG AGCTGGTAGG AGCCGAGTGC CACAGGCATT CTGCATTGCT 840

45

CTACTCTTAG GTTTGTGTGT GTGATCCTTC CCTCCCTGT CCGCCACTCC TCCCTCCTCT 900

GGCTATCCTA CCCTGTCTGT GGGCTCTTTT ACTACCAGCC TATCTGTGCG GACTGTCATG 960

GCATTTAGTT CAGAGTGGAN GGGCTTTGGS CTGAAATAAA ATGCAAGTAT TTAAAAAAAA 1020

50

AAAAAAAAA A 1031

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(2) INFORMATION FOR SEQ ID NO: 71:

(i) SEQUENCE CHARACTERISTICS:

60

- (A) LENGTH: 855 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 71:

5	AGCTATTGAC ACTTCCTGGT GGGATCCGAG TGAGGCGACG GGGTAGGGGT TGGCGCTCAG	60
	CGGGCGACCA TGGCGTATCA CGGCCTCACT GTGCCTCTCA TTGTGATGAG CGTGTCTCTGG	120
	GGCTTCGTCG GCTTCTTGGT GCCTTGGTTC ATCCCTAAGG GTCCTAACCG GGGAGTTATC	180
10	ATTACCATGT TGGTGACCTG TTCAGTTTGC TGCTATCTCT TTTGGCTGAT TGCAATTCTG	240
	GCCCAACTCA ACCCTCTCTT TGGACCGCAA TTGAAAAATG AAACCATCTG GTATCTGAAG	300
15	TATCATTGGC CTTGAGGAAG AAGACATGCT CTACAGTGCT CAGTCTTTGA GGTCACGAGA	360
	AGAGAAATGCC TTCTAGATGC AAAATCACCT CCAAACCAGA CCACTTTTCT TGA CTTCGCCT	420
	GTTTTGGCCA TTAGCTGCCT TAAACGTTAA CAGCACATTT GAATGCCTTA TTCTACAATG	480
20	CAGCGTGTTC TCCTTTCCT TTTTGGCACT TTGGTGAATT ACGTGCCTCC ATAACCTGAA	540
	CTGTGCCGAC TCCACAAAAC GATTATGTAC TCTTCTGAGA TAGAAGATGC TGTCTCTCTG	600
25	AGAGATACGT TACTCTCTCC TTGGAATCTG TGGATTTGAA GATGGCTCCT GCCTTCTCAC	660
	GTGGGAATCA GTGAAGTGTT TAGAACTGC TGCAAGACAA ACAAGACTCC AGTGGGGTGG	720
	TCAGTAGGAG AGCACGTTCA GAGGGAAGAG CCATCTCAAC AGAATCGCAC CAACTATAC	780
30	TTTCAGGATG AATTCTTCT TTCTGCCATC TTTTGAATA AATATTTTCC TCCTTTCTAW	840
	RRAAAAAAA ANANN	855

35

(2) INFORMATION FOR SEQ ID NO: 72:

40	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 1274 base pairs
	(B) TYPE: nucleic acid
	(C) STRANDEDNESS: double
	(D) TOPOLOGY: linear

45

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:

	GGCAGAGCTT AGAGTGTTGA AAAGGCAACC AGGTTGGCCG TAAGTGCCTG CTGGAATGCG	60
50	TGTGCCTCCA CACGGGTCTG GGCATCCGGA CTGATAACCA GCCGGCCAGA CTGAGGGATG	120
	GAAGGCACTG AGATGGGGGC CCGTCCAGGC GGACACCCGC AGAAATGGAG CTTTCTGTGG	180
	TCTCTTGCAC TCTGGCTGCC TCTTGCCCTC TCTGTGTCTC TCTTCTTTGG TCTCTCCCTC	240
55	TCTCCTCCTC AGCCTGGTCT TTCTCTTTGG TGCACACTTA GTTATTGTTG TGAGCAATGG	300
	AAGTTCAAAG GAACTCCCTC TCCAGCTCTT CTGAATCTTG GGACACAGCC TAAAAAGGAC	360
60	AAAAAGTTAG AAGACAGCAT AGCAACTCAG CTCAGGGAGC TACACAGAGAA AAATAGCAAC	420

	TGATGTGGGT GCTTTTTTTT TTTTTTTAAT TTGAATAAAA AGAATTAGAA GTGATGTCCT	480
5	TTTATAAAAT GCCTTCTCCC CCTTCCCGCC TACAGTCTCT TCCTCTCCCC TTAGAGGGGG	540
	GAAAGTGTAT AAACCTACAG GGTGTGTAGT CTGAAAAGAG GATCCCCCTC ACCCCCACCC	600
	TGGGCAGAGC AGTGGGGGTT GGGGGGTGGG AGAGGGGGAC ACAGATCCTG GCACACTGTG	660
10	GATATTTCTT GCAGATTGCA GTCTCTTGTG GCCCAAACAG GTTAGGTAGA CTATCGCCTC	720
	TGGCAGGTGC CACCTTTTGG TACCAACATG TTCTGAGGTG TTAGGATTTG GGTGGGTTT	780
15	TTTTTGTTG TTTTTTTTTT CCTTTTGGTC TTTTTTTTTT TCTCCTTTTA AAGAAAAGCT	840
	AAAGGCCGCT GTGAGTCTTG GTGGCAGGCT CTCATGGAT GTAGCATATC GAAGATAATT	900
	TTTATACTGC ATTTTATGG ATTATTTTGT AATGTGTGAT TCCGTCTGCT GAGGAGGTGG	960
20	GAGGGGCTCC AGGGAAGCC ACCCACCTTC AGTGAGGTG CTCCCCAGCT GAGCGCACCG	1020
	GGCATGGGAT GTGAGGCTG GCGACACACC CTGTGCCTCT CCAAGGCTGG GCGCGTGGG	1080
25	CGTCCAGAGT CTCTCTGGGT CTCAGATGTC CATCTGCCAC CTCTTGTTAA GGCTCTAGCC	1140
	AGAAGGGAGG GTGAGGGTAG AAGAAAGTTA TTCCCGAAGA AAAAAAGAAT GAAAAGTCAT	1200
	TGTACTGAAC TGTTTTATA TTTTAAAAG TTACTATTWA AAGGTAAAAA AAAGGGGGG	1260
30	CCCGGTACCC AATT	1274

35 (2) INFORMATION FOR SEQ ID NO: 73:

(i) SEQUENCE CHARACTERISTICS:

- 40 (A) LENGTH: 688 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:

45	GGCAGGAGTG GAGGCAATGC CAGCTCCAGG ACAGAGGCTC AGGTGCCCAA CGGGCAAGGC	60
	AGCCCAGGGG GCTGTGTCTG TTCAAGTCAG GCTTCCCCGG CCCTCGCGCA CAGCGCTTCC	120
50	ACGGGCAGCC CGGGGCCCCA CCCCACGCAC TGAAGAGGCC GCCTGGGCTG CCATGGCCCT	180
	GACCTTCCTG CTGGTGCTGC TCACCTGGC CACGTCTGCA CACGGCTGCA CAGAACTTC	240
	CGACGCGGGG AGAGCATCTA CTGGGGGCCC ACAGCGGACA GCGAGGACAC AGTGGCTGCT	300
55	GTGCTGAAGC GGAGGTGCT GCAGCCCTCG CGCCGGGTCA AGCGTTCGGG CCGGAGACCC	360
	CTCTCCCGCC CACGCCGAC AGCGGCCCGG AAGGCGAGAG CTCTGAGTGA CCGCCTGGGA	420
60	CCTGCCACTG TGGCGTGCGG CTCCTCCCCG CGCCGCGAGG CCGCGACCTC TGCCACGTGG	480

ACCGCGCGCG GGGCGCTCCC TGGTGGCGAT GGC GCGGCAC TGGCCGAGCA CTGCGGGGGC 540
 TTTCTCTCTT GTTGGTTGCT GAGTGGGCGG CCAAGGGGAG AAAAGGAGCC GCTTCTGCCT 600
 5 CCCTTGCCAA AACTCCGTTT CTAATTAAT TATTTTAGT AGAAAAAAAA AAAAAAAAAA 660
 AAAAAAAAAA AAAAAAAAAA AAAAAAA 688

10

(2) INFORMATION FOR SEQ ID NO: 74:

15 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1890 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:

GAGCAGGAGA GAAGGCACCG CCCACCCCG CCTCCAAAGC TAACCCTCGG GCTTGAGGGG 60
 AAGAGGCTGA CTGTACGTTT CTTCTACTCT GGCACCACTC TCCAGGCTGC CATGGGGCCC 120
 25 AGCACCCCTC TCCTCATCTT GTTCCTTTTG TCATGGTCGG GACCCCTCCA AGGACAGCAG 180
 CACCACCTTG TGGAGTACAT GGAACGCCGA CTAGTGCTT TAGAGGAACG GCTGGCCCAG 240
 30 TGCCAGGACC AGAGTAGTCG GCATGCTGCT GAGCTGCGGG ACTTCAAGAA CAAGATGCTG 300
 CCACTGCTGG AGGTGGCAGA GAAGGAGCGG GAGGCACTCA GAACTGAGGC CGACACCATC 360
 TCCGGGAGAG TGGATCGTCT GGAGCGGGAG GTAGACTATC TGGAGACCCA GAACCCAGCT 420
 35 CTGCCCTGTG TAGAGTTTGA TGAGAAGGTG ACTGGAGGCC CTGGGACCAA AGGCAAGGGA 480
 AGAAGGAATG AGAAGTACGA TATGGTGACA GACTGTGGCT ACACAATCTC TCAAGTGAGA 540
 40 TCAATGAAGA TTCTGAAGCG ATTTGGTGGC CCAGCTGGTC TATGGACCAA GGATCCACTG 600
 GGGCAAACAG AGAAGATCTA CGTGTTAGAT GGGACACAGA ATGACACAGC CTTTGTCTTC 660
 CCAAGGCTGC GTGACTTCAC CCTTGCCATG GCTGCCCGGA AAGCTTCCCG AGTCCGGGTG 720
 45 CCCTTCCCTT GGTAGGCAC AGGGCAGCTG GTATATGGTG GCTTCTTTTA TTTTGCTCGG 780
 AGGCCTCCTG GAAGACCTGG TGGAGGTGGT GAGATGGAGA ACACTTTGCA GCTAATCAAA 840
 50 TTCCACCTGG CAAACCGAAC AGTGGTGGAC AGCTCAGTAT TCCAGCAGA GGGGCTGATC 900
 CCCCCTACG GCTTGACAGC AGACACCTAC ATCGACCTGG CAGCTGATGA GGAAGGTCTT 960
 TGGGCTGTCT ATGCCACCCG GGAGGATGAC AGGCACTTGT GTCTGGCCAA GTTAGATCCA 1020
 55 CAGACACTGG ACACAGAGCA GCAGTGGGAC ACACCATGTC CCAGAGAGAA TGCTGAGGCT 1080
 GCCTTTGTCA TCTGTGGGAC CCTCTATGTC GTCTATAACA CCGCTCCTGC CAGTCGGGCC 1140
 60 CGCATCCAGT GCTCCTTTGA TGCCAGCGGA CCCTGACCCC TGAACGGGCA GCACTCCCTT 1200

ATTTTCCCCG CAGATATGGT GCCCATGCCA GCCTCCGCTA TAACCCCCGA GAACGCCAGC 1260
 TCTATGCCTG GGATGATGGC TACCAGATTG TCTATAAGCT GGAGATGAGG AAGAAAGAGG 1320
 5 AGGAGGTTTG AGGAGCTAGC CTTGTTTTTT GCATCTTTCT CACTCCCATA CATTTATATT 1380
 ATATCCCCAC TAAATTTCTT GTTCCTCATT CTTCAAATGT GGGCCAGTTG TGGCTCAAAT 1440
 10 CCTCTATATT TTTAGCCAAT GGCAATCAAA TTCTTTCAGC TCCTTTGTTT CATACGGAAC 1500
 TCCAGATCCT GAGTAATCCT TTTAGAGCCC GAAGAGTCAA AACCTCAAT GTTCCCTCCT 1560
 GCTCTCCTGC CCCATGTCAA CAAATTTAGC GCTAAGGATG CCCCAGACCC AGGGCTCTAA 1620
 15 CCTTGTATGC GGGCAGGCCC AGGGAGCAGG CAGCAGTGTT CTTCCCCTCA GAGTGACTTG 1680
 GGGAGGGAGA AATAGGAGGA GACGTCCAGC TCTGTCCTCT CTTCTCACT CCTCCCTTCA 1740
 20 GTGTCCTGAG GAACAGGACT TTCTCCACAT TGTTTTGTAT TGCAACATTT TGCATTAAAA 1800
 GGAAATCCA CTGCAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAACGG CACGAGGGGG 1860
 GGTCCCGTAC CCAATNGCCC TCACATGCAT 1890
 25

(2) INFORMATION FOR SEQ ID NO: 75:
 30 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1133 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 35 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 75:

GCCGGTCTGA GTGCAGAGCT GCTGTCATGG CGGCCGCTCT GTGGGGCTTC TTTCCCGTCC 60
 40 TGCTGCTGCT GCTGCTATCG GGGGATGTCC AGAGCTCGGA GGTGCCCCGG GCTGCTGCTG 120
 AGGGATCGGG AGGGAGTGGG GTCGGCATAG GAGATCGCTT CAAGATTGAG GGGCGTGCAG 180
 45 TTGTTCCAGG GGTGAAGCCT CAGGACTGGA TCTCGGCGGC CCGAGTGCTG GTAGACGGAG 240
 AAGAGCACGT CGGTTTCCTT AAGACAGATG GGAGTTTGT GTTCATGAT ATACCTTCTG 300
 GATCTTATGT AGTGGAAGTT GTATCTCCAG CTTACAGATT TGATCCCGTT CGAGTGGATA 360
 50 TCACTTCGAA AGGAAAAATG AGAGCAAGAT ATGTGAATTA CATCAAAACA TCAGAGGTTG 420
 TCAGACTGCC CTATCCTCTC CAAATGAAAT CTTCAGGTCC ACCTTCTTAC TTTATTAAAA 480
 55 GGAATCGTG GGGCTGGACA GACTTCTTAA TGAACCAAT GGTATGATG ATGGTTCTTC 540
 CTTTATTGAT ATTTGTGCTT CTGCCTAAAG TGGTCAACAC AAGTGATCCT GACATGAGAC 600
 GGGAAATGGA GCAGTCAATG AATATGCTGA ATTCCAACCA TGAGTTCCCT GATGTTTCTG 660
 60

AGTTCATGAC AAGACTCTTC TCTTCAAAT CATCTGGCAA ATCTAGCAGC GGCAGCAGTA 720
 AAACAGGCAA AAGTGGGGCT GGCAAAGGA GGTAGTCAGG CCGTCCAGAG CTGGCATTG 780
 5 CACAAACACG GCAACACTGG GTGGCATCCA AGTCTTGGA AACC GTGTGA AGCAACTACT 840
 ATAAACTTGA GTCATCCCGA CGTTGATCTC TTACAACTGT GTATGTTAAC TTTT TAGCAC 900
 10 ATGTTTTGTA CTTGGTACAC GAGAAAACCC AGCTTTCATC TTTTGTCTGT ATGAGGTCAA 960
 TATTGATGTC ACTGAATTAA TTACAGTGTC CTATAGAAAA TGCCATTAAAT AAATTATATG 1020
 AACTACTATA CATTATGTAT ATTAATTAAA ACATCTTAAT CCAGAAAAAA AAAAAAARAA 1080
 15 AACTCGAGGG GGGGCCCGGT ACCCAATTTN CCAAATGGGA GTCGTAAAA ATC 1133

20 (2) INFORMATION FOR SEQ ID NO: 76:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 585 base pairs
 (B) TYPE: nucleic acid
 25 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:

30 ATGTTTACAA TGTTGTGTAT AAATGGGACA ACTCCTCGCC CTCTACCTGT CCCCTCCCC 60
 TTTGGTTGTA TGATTTTCTT CTTT TTAAG AACCCCTGGA AGCAGCGCCT CCTTCAGGGT 120
 TGGCTGGGAG CTCGGCCCAT CCACCTCTTG GGGTACCTGC CTCTCTCTCT CCTGTGGTGT 180
 35 CCCTTCCCTC TCCCATGTGC TCGGTGTTCA GTGGTGTATA TTTCTTCTCC CAGACATGGG 240
 GCACACGCCC CAAGGGACAT GATCCTCTCC TTAGTCTTAG CTCATGGGGC TCTTTATAAG 300
 40 GAGTTGGGGG GTAGAGGCAG GAAATGGGAA CCGAGCTGAA GCAGAGGCTG AGTTAGGGGG 360
 CTAGAGGACA GTGCTCCTGG CCACCCAGCC TCTGCTGAGA ACCATTCCTG GGATTAGAGC 420
 TGCCTTTCCC AGGGAAAAAG TGTCGTCTCC CCGACCCTCC CGTGGGCCCT GTGGTGTGAT 480
 45 GCTGTGTCTG TATATTCTAT ACAAAGGTAC TTGTCCTTTC CCTTTGTAAA CTACATTGA 540
 CATGGATTAA ACCAGTATAA ACAGTTAAAA AAAAAAAAAA AAAAA 585

50

(2) INFORMATION FOR SEQ ID NO: 77:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 577 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 60 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:

	GGCACGAGGC CTTCGAGAAC TTCTACTTGC CTGCCTCCCT GCCTCTGGCC ATGGCCTGCC	60
5	GGTGCCTCAG CTTCTTCTG ATGGGGACCT TCCTGTCACT TTCCCAGACA GTCCTGGCCC	120
	AGCTGGATGC ACTGCTGGTC TTCCCAGGCC AAGTGGCTCA ACTCTCCTGC ACGCTCAGCC	180
10	CCCAGCACGT CACCATCAGG GACTACGGTG TGTCTTGTA CCAGCAGCGG GCAGGCAGTG	240
	CCCCTCGATA TCTCTCTAC TACCGCTCGG AGGAGGATCA CCACCGGCCT GCTGACATCC	300
	CCGATCGATT CTCGGCAGCC AAGGATGAGG CCCACAATGC CTGTGTCTC ACCATTAGTC	360
15	CCGTGCAGCC TGAAGACGAC GCGGATTACT ACTGCTCTGT TGGCTACGGC TTTAGTCCCT	420
	AGGGGTGGGG TGTGAGATGG GTGCCTCCCC TCTGCCTCCC ATTTCTGCCC CTGACCTTGG	480
20	GTCCCTTTTA AACTTTCTCT GAGCCTTGCT TCCCTCTGT AAAATGGGTT AATAATATTC	540
	AACATGTCAA CAACAAAAA NAAAAWAAA AACTCGA	577

25

(2) INFORMATION FOR SEQ ID NO: 78:

(i) SEQUENCE CHARACTERISTICS:

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- (A) LENGTH: 2278 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:

35

	GTAATTCGGC ACGAGGCGCC CAACATGGCG GGTGGGCGCT GCGGCCCGCA SCTAACGGCG	60
	CTCCTGGCCG CCGGATCGC GGCTGTGGCG GCGACGGCAG GCCCGAGGA GGCCGCGCTG	120
40	CCGCCGAGC AGAGCCGGT CCAGCCCATG ACCGCCTCA ACTGGACGCT GGTGATGGAG	180
	GGCGAGTGGA TGCTGAAAT TTACGCCCCA TGGTGTCCAT CCTGCCAGCA GACTGATTCA	240
45	GAATGGGAGG CTTTTCGCAA GAATGGTGAA ATACTTCAGA TCAGTGTGGG GAAGGTAGAT	300
	GTCATTCAAG AACCAGGTTT GAGTGGCCGC TTCTTTGTCA CCACTCTCCC AGCATTTTPT	360
	CATGCAAAGG ATGGGATATT CCGCCGTAT CGTGGCCAG GAATCTTCGA AGACCTGCAG	420
50	AATTATATCT TAGAGAAGAA ATGGCAATCA GTCGAGCCTC TGAATGGCTG GAAATCCCCG	480
	GCTTCTCTAA CGATGTCTGG AATGGCTGGT CTTTTCAGCA TCTCTGGCAA GATATGGCAT	540
55	CTTCACAACT ATTTACAGT GACTCTTGA ATTCCTGCTT GGTGTCTTA TGTCTTTTTC	600
	GTCATAGCCA CCTTGGTTTT TGGCCTTTTT ATGGGTCTGG TCTTGGTGGT AATATCAGAA	660
	TGTTTCTATG TGCCACTTCC AAGGCATTTA TCTGAGCGTT CTGAGCAGAA TCGGAGATCA	720
60	GAGGAGGCTC ATAGAGCTGA ACAGTTGCAG GATGCGGAGG AGGAAAAAGA TGATTCAAAT	780

	GAAGAAGAAA ACAAAGACAG CCTTG TAGAT GATGAAGAAG AGAAAGAAGA TCTTGCGAT	840
	GAGGATGAAG CAGAGGAAGA AGAGGAGGAG GACAACTTGG CTGCTGGTGT GGATGAGGAG	900
5	AGAAGTGAGG CCAATGATCA GGGGCCCCCA GGAGAGGACG GTGTGACCCG GGAGGNAAGT	960
	AGAGCCTGAG GAGGCTGAAG AAGGCATCTC TGAGCAACCC TGCCCAGCTG ACACAGAGGT	1020
10	GGTGGAAGAC TCCTTGAGGC AGCGTAAAAG TCAGCATGCT GNCAAGGGAC TG TAGATTTA	1080
	ATGATGCGTT TTCAAGAATA CACACCAAAA CAATATGTCA GCTTCCCTTT GGCCTGCAGT	1140
	TTGTACCAAA TCCTTAATTT TTCCTGAATG AGCAAGCTTC TCTTAAAAGA TGCTCTCTAG	1200
15	TCATTTGGTC TCATGGCAGT AAGCCTCATG TATACTAAGG AGAGTCTTCC AGGTGTGACA	1260
	ATCAGGATAT AGAAAAACAA ACGTAGTGTN TGGGATCTGT TTGGAGACTG GGATGGGAAC	1320
20	AAGTTCATTT ACTTAGGGGT CAGAGAGTCT CGACCAGAGG AGGCCATTCC CAGTCCTAAT	1380
	CAGCACCTTC CAGAGACAAG GCTGCAGGCC CTGTGAAATG AAAGCCAAGC AGGAGCCTTG	1440
	GNCTCTAGGC ATCCCCAAAG TGTAACGTAG AAGCCTTGCA TCCTTTTCTT GTGTAAAGTA	1500
25	TTTATTTTGT TCAAATTGCA GGAAACATCA GGCACCACAG TGCATGAAAA ATCTTTCACA	1560
	GCTAGAAATT GAAAGGGCCT TGGGTATAGA GAGCAGCTCA GAAGTCATCC CAGCCCTCTG	1620
30	AATCTCCTGT GCTATGTTTT ATTTCTTACC TTTAATTTTT CCAGCATTTC CACCATGGGC	1680
	ATTCAGGCTC TCCACACTCT TCACTATTAT CTCTTGGTCA GAGGACTCCA ATAACAGCCA	1740
	GGTTTACATG AACTGTGTTT GTTCATTCTG ACCTAAGGGG TTTAGATAAT CAGTAACCAT	1800
35	AACCCCTGAA GCTGTGACTG CCAAACATCT CAAATGAAAT GTTGTGGCCA TCAGAGACTC	1860
	AAAAGGAAGT AAGGATTTTA CAAGACAGAT TAAAAAATAA TTGTTTTGTC CAAAATATAG	1920
40	TTGTTGTTGA TTTTTTTTAA AGTTTTCTAA GCAATATTTT TCAAGCCAGA AGTCCTCTAA	1980
	GTCTTGCCAG TACAAGGTAG TCTGTGAAG AAAAGTTGAA TACTGTTTTG TTTTCATCTC	2040
	AAGGGGTTC CTGGGTCTTG AACTACTTTA ATAATACTA AAAAACCCT TCTGATTTTC	2100
45	CTTCAGTGAT GTGCTTTTGG TGAAAGAATT AATGAACTCC AGTACCTGAA AGTGAAAGAT	2160
	TTGATTTTGT TTCCATCTTC TGTAATCTTC CAAAGAATTA TATCTTTGTA AATCTCTCAA	2220
50	TACTCAATCT ACTGTAAGTA CCCAGGGAGG CTAATTTCTT TAAAAAATAA AAAAAA	2278

55 (2) INFORMATION FOR SEQ ID NO: 79:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1143 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

60

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:

5 CCCCTCCAAC TCTCAACCCA CTTCTCCAGC CAGCGCCCCA GCCCTCCCGC CGCCCGCTCG 60
 CAGGTCCCGA GGAGCGCAGA CTGTGTCCCT GACAATGGGA ACAGCCGACA GTGATGAGAT 120
 GGCCCCGGAG GCCCCACAGC ACACCCACAT CGATGTGCAC ATCCACCAGG AGTCTGCCCT 180
 10 GGCCAAGCTC CTGCTCACCT GCTGCTCTGC GCTGCGGCCC CGGGCCACCC AGGCCAGGGG 240
 CAGCAGCCGG CTGCTGGTGG CCTCGTGGGT GATGCAGATC GTGCTGGGGA TCTTGAGTGC 300
 15 AGTCCTAGGA GGATTTTTCT ACATCCGCGA CTACACCCTC CTCGTACCTC CGGGAGCTGC 360
 CATCTGGACA GGGGCTGTGG CTGTGCTGGC TGGAGCTGCT GCCTTCATTT ACGAGAAACG 420
 GGGTGGTACA TACTGGGCCC TGCTGAGGAC TCTGCTAGCG CTGGCAGCTT TCTCCACAGC 480
 20 CATCGCTGCC CTCAAACCTT GGAATGAAGA TTTCCGATAT GGCTACTCTT ATTACAACAG 540
 TGCCTGCCGC ATCTCCAGCT CGAGTGACTG GAACACTCCA GCCCCACTC AGAGTCCAGA 600
 25 AGAAGTCAGA AGGCTACACC TATGTACCTC CTTCATGGAC ATGCTGAAGG CCTTGTTTAC 660
 AACCCTTCAG GCCATGCTCT TGGGTGTCTG GATTCTGCTG CTTCTGGCAT CTCTGGCCCC 720
 TCTGTGGCTG TACTGCTGGA GAATGTTCCC AACCAAAGG AAAAGAGACC AGAAGGAAAT 780
 30 GTTGGAAGTG AGTGAATCT AGCCATGCCT CTCCTGATTA TTAGTGCCTG GTGCTTCTGC 840
 ACCGGGCGTC CCTGCATCTG ACTGCTGGAA GAAGAACCAG ACTGAGGAAA AGAGGCTCTT 900
 35 CAACAGCCCC AGTTATCCTG GCCCCATGAC CGTGGCCACA GCCCTGCTCC AGCAGCACTT 960
 GCCCATTCCT TACACCCCTT CCCCATCCTG CTCGCTTCA TGTCCCCTCC TGAGTAGTCA 1020
 TGTGATAATA AACTCTCATG TTATTGTTNN NAAAAAAAAA AAAAAAAAAA AATTGGGGG 1080
 40 GGGGCCGGTA CCCATTGGGC CTNNGGGGN GGTTTAAAT TAATGGGGG GGTTTAAAG 1140
 GGN 1143
 45

(2) INFORMATION FOR SEQ ID NO: 80:

50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 557 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:

GGCAGAGAGC AGATGGCCTT GACACCAGCA GGGTGACATC GGTATTCCT ACTTCTCTGC 60
 60 TCCCCACAG TTCCTCTGGA CTTCTCTGGA CCACAGTCCT CTGCCAGACC CCTGCCAGAC 120

5 CCCAGTCCAC CATGATCCAT CTGGGTCACA TCCTCTTCCT GCTTTTGCTC CCAGTGGCTG 180
 CAGCTCAGAC GACTCCAGGA GAGAGATCAT CACTCCCTGC CTTTACCCT GGCACCTCAG 240
 GCTCTTGTTT CGGATGTGGG TCCCTCTCTC TGCCGCTCCT GGCAGGCCTC GTGGCTGCTG 300
 ATGCGGTGGC ATCGCTGCTC ATCGTGGGG CGGTGTTCTT GTGCGCACGC CCACGCCGCA 360
 10 GCCCCGCCCA AGAAGATGGC AAAGTCTACA TCAACATGCC AGGCAGGGGC TGACCCTCCT 420
 GCAGCTTGA CTTTGACTT CTGACCCTCT CATCCTGGAT GGTGTGTGGT GGCACAGGAA 480
 CCCCCGCCCC AACTTTTGA TTGTAATAAA ACAATTGAAA CACCAAAAAA AAAAAAAAAA 540
 15 AAAAAAAAAA AANTCGA 557

20

(2) INFORMATION FOR SEQ ID NO: 81:

(i) SEQUENCE CHARACTERISTICS:

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- (A) LENGTH: 795 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:

30

GCCCGGGCGA TGTGGAGCGC GGGCCGCGGC GGGGCTGCCT GGCCGGTGCT GTTGGGGCTG 60
 CTGCTGGCGC TGTAGTGCC GGGCGGTGGT GCCGCCAAGA CCGGTGCGGA CTCGTGACCT 120
 35 GCGGGTCGGT GCTGAAGCTG CTCAATACGC ACCACCGCGT GCGCTGCACT CGCACGACAT 180
 CAAATACGGA TCCGGCAGCG GCCAGCAATC GGTGACCGGC GTAGAGGCGT CGGACGACGC 240
 MAATAGCTAC TGGCGGATCC GCGGCGGCTC GGAGGGCGGG TCCCCGCGCG GTCCCCGGT 300
 40 GCGCTGCGGG CAGGCGGTGA GGCTCACGCA TGTSCCTTACG GGCAAGAACY TGCACACGCA 360
 CCAYTTCCCG TCGCCGCTGT CCAACAACCA GGAGGTGAGT GCTTTTGGGG AAGACGGCGA 420
 45 GGGCGACGAC CTGGACCTAT GGACAGTGCG CTGCTCTGGA GAGCACTGGG AGCGTGAGGC 480
 TGCTGTGCCT TCCAGCATGT GGGCACCTCT GTGTTCTCTT TACTCAGCGG TGAGCAGTAT 540
 GGAAGCCCCA TCCGTGGGCA GCATGAGGTC CACGGCATGT GAGTTGCGAA CACGCACAAT 600
 50 ACGTGGAAGG CCATGGAAGG CATCTTCATC AAGCCTAGT TCGAGTCTCT TGCAGGTCAC 660
 GATGAACTCT GAGTGTGTGG ATGGATGGGT GGATGGAGG TCTTGGTTCT GCGCTCTGCA 720
 55 GGGCCACTCT TGGCAGAGAC TTTGGGTTTG TAGGGTCTCT TTTTCTTT TTTGATTAA 780
 GAATGTTGGT CTATG 795

60

(2) INFORMATION FOR SEQ ID NO: 82:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 1324 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 82:

NAGGCTTTAA AGCGCTACC CTGCCTGCAG GTGAGCACTG GTGTGTGAGA GCCAGGCGTC 60
 CCTCTGCCTG CCCACTCAGT GGCAACACCC GGGAGCTGTT TTGTCTTTTG TGGAGCCTCA 120
 15 GCAGTTCCCT CTTTCAGAAC TCACTGCCAA GAGCCCTGAA CAGGAGCCAC CATGCAGTGC 180
 TTCAGCTTCA TTAAGACCAT GATGATCCTC TTCAATTGTC TCATCTTTCT GTGTGGTGCA 240
 20 GCCCTGTTGG CAGTGGGCAT CTGGGTGTCA ATCGATGGGG CATCCTTTCT GAAGATCTTC 300
 GGGCCACTGT CGTCCAGTGC CATGCAGTTT GTCAACGTGG GCTACTTCCT CATCGCAGCC 360
 GCGTGTGTGG TCTTTGCTCT TGGTTTCCTG GGCTGCTATG GTGCTAAGAC TGAGAGCAAG 420
 25 TGTGCCCTCG TGACGTTCCT CTCATCCTC CTCCTCATCT TCATTGCTGA GGTTCAGCT 480
 GCTGTGGTCG CCTTGGTGTA CACCACAATG GCTGAGCACT TCCTGACGTT GCTGGTAGTG 540
 30 CCTGCCATCA AGAAAGATTA TGGTTCCTAG GAAGACTTCA CTCAAGTGTG GAACACNACC 600
 ATGAAAGGGC TCAAGTGCTG TGGCTTCACC AACTATACGG ATTTTGAGGA CTCACCTAC 660
 TTCAAAGAGA ACAGTGCCTT TCCCCATTC TGTTGCAATG ACAACGTAC CAACACAGCC 720
 35 AATGAAACCT GCACCAAGCA AAAGGCTCAC GACCAAAAAG TAGAGGGTTG CTTCAATCAG 780
 CTTTGTATG ACATCCGAAC TAATGCAGTC ACCGTGGGTG GTGTGGCAGC TGGAAATGGG 840
 40 GGCTCGAGC TGGCTGCCAT GATTGKTCC ATGTATCTGT ACTGCAATCT ACAATAAGTC 900
 CACTTCTGCC TCTGCCACTA CTGCTGCCAC ATGGGAAC TGGAAGAGGCA CCCTGGCAAG 960
 CAGCAGTGAT TGGGGGAGGG GACAGGATCT AACAAATGCA CTGGGGCCAG AATGGACCTG 1020
 45 CCCTTCTGTC TCCAGACTTG GGGCTAGATA GGGACCACTC CTTTAGGCA TGCCTGACTT 1080
 TCCTTCCATT GGTGGGTGGA TGGGTGGGGG GCATTCCAGA GCTTTAAGG TAGCCAGTTC 1140
 50 TGTGCCCCAT TCCCCAGTC TATTAAACCC TTGATATGCC GCTTAAGCT AGTGGTGATC 1200
 CCAGTGCTCT ACTGGGGGAT GAGAGAAAGG CATTTTATAG GCTTATATA AGTGAAATCA 1260
 GCAGAGCCTC TGGGTGGATG TGTAGAAGGC ACTTCAAATC TATTAATCT TGTACAAATG 1320
 55 TAAA 1324

(2) INFORMATION FOR SEQ ID NO: 83:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 1494 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83:

10 CTCAGGCTTC TGTCTCACTT TTCCGGGGGG GGGATTAGGG CAAGGAGGGC ATGAGGGACT 60
 GTCTCTCCCT AAAACCCAGA CCCCTGTTC CCACTCAGTT CTTCTTCATC CTCCTCCTCA 120
 15 TCTTCATTGC TGAGGTTGCA GCTGCTGTGG TCGCCTTGGT GTACACCACA ATGGTGAGAC 180
 ACTGGGATGG AGGAAGGGAA GAAGATTGGG CAAAACCCTG GGAGTGGGCT GTGGCCTGTG 240
 20 AATGGCCACC TTCTGTACCA GCCCCTAAAC ACTGGCCTGC CTCACCCAGG CTGAGCACTT 300
 CCTGACGTTG CTGGTAGTGC CTGCCATCAA GAAAGATTAT GGTTCACAGG AAGACTTCAC 360
 TCAAGTGTGG AACACCACCA TGAAAGGGGT AAGGTTGGCT GGGGAGGTT TTAGGGTGA 420
 25 GAGAAAGAAG CAAGCCCCA CCTCCACCCT CATCTTGTCT CCAGCTCAAG TGCTGTGGCT 480
 TCACCAACTA TACGGATTTT GAGGACTCAC CCTACTTCAA AGAGAACAGT GCCTTTCCCC 540
 CATCTGTGTT CAATGACAAC GTCACCCAAC ACAGCCCAAT GAAACCTGCA CCAAGCAAAA 600
 30 GGCTCACASAC CNAAAARTAN AGGTGTGGGC TGGCATGAGT GGGTGGGAC TGTTTTCATG 660
 GCCTCAGAGT GGCAAACGGG GATGGGAGTA GGCAGCTGC CAACTATAAA TGCTCTTTTC 720
 35 TCTTCCYGAA GGGTTGCTTC AATCAGCTTT TGTATGACAT CCGAACTAAT GCAGTCACCG 780
 TGGGTGGTGT GGCAGCTGGA ATTGGGGGCC TCGAGGTAAG CAGATSAGGA GCTGGGACTG 840
 GGACATGGGC ATGAGACCAG GGCTGCTCAA CCCATCTGAG GCCTCTCTGG AGGAAACAGA 900
 40 CTTCTAACTG GGCCTCAGGT AGGGTGTCTG TGGGACAGGC TTCAGGATCC CTATCATGTT 960
 CCCTCATCTC TCCCTGTTCC TCCCTCTCCA GCTGGCTGCC ATGATTGTGT CCATGTATCT 1020
 45 GTACTGCAAT CTACAATAAG TCCACTTCTG CCTCTGCCAC TACTGCTGCC ACATGGGAAC 1080
 TGTGAAGAGG CACCCTGGCA AGCAGCAGTG ATTGGGGGAG GGGACAGGAT CTAACAATGT 1140
 CACTTGGGCC AGAATGGACC TGCCCTTTCT GCTCCAGACT TGGGGCTAGA TAGGGACCAC 1200
 50 TCCTTTTAGC GATGCCTGAC TTTCCTTCCA TTGGTGGGTG GATGGGTGGG GGGCATTCCA 1260
 GAGCCTCTAA GGTAGCCAGT TCTGTTGCC ATTCCCCAG TCTATTAAAC CCTTGATATG 1320
 55 CCCCCTAGGC CTAGTGGTGA TCCCAGTGCT CTACTGGGGG ATGAGAGAAA GGCATTTTAT 1380
 AGCCTGGGCA TAAGTGAAAT CAGCAGAGCC TCTGGGTGGA TGTGTAGAAG GCACTTCAAA 1440
 60 ATGCATAAAC CTGTTACAAT GTTAAAAAAA AAAAAAAAAA AACTCGACTC TGCC 1494

(2) INFORMATION FOR SEQ ID NO: 84:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1285 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:

5	GCTACGTGGC TGGCATGCAT GGGAAACGAGG CCCTGGGGCG GGAGTTGCTT CTGCTCCTGA	60
15	TGCAGTTTCCT GTGCCATGAG TTCCTGCGAG SGAACCCACG GGTGACCCGG CTGCTCTCTG	120
	AGATGCGCAT TCACCTGCTG CCCTCCATGA ACCCTGATGG CTATGAGATC GCCTACCACC	180
20	GGGGTTCAGA RCTGGTGGGC TGGGCCGARG GCCGCTGGAA CAACCAGAGC ATCGATCTTA	240
	ACCATAATTT TGCTGAMCTC AACACACCAC TGTGGGAAGC ACAGGACGAT GGAAGGTGC	300
25	CCCACATCGT CCCCACCAT CACCTGCCAT TGCCCACTTA CTACACCTG CCAATGCCA	360
	CCGTGGCTCC TGAAACGCGG GCAGTAATCA AGTGGATGAA GCGGATCCCC TTTGTGCTAA	420
	GTGCCAACCT CCACGGGGT GAGCTCGTGG TGTCCTACCC ATTCGACATG ACTCGCACCC	480
30	CGTGGGCTGC CCGCGAGCTC ACGCCACAC CAGATGATGC TGTGTTTCGC TGGCTCAGCA	540
	CTGTCTATGC TGGCAGTAAT CTGGCCATGC AGGACACCAG CCGCCGACCC TGCCACAGCC	600
35	AGGACTTCTC CGTGCACGGC AACATCATCA ACGGGGCYTG ACTNGGCACA CGGTCCCCGG	660
	GANGCATGAA TGAYTTCAGC TACCTACACA CCAACTGCTT TGAGGTCAC TGGGAGCTGT	720
	SCTGTGACAA GTTCCTCAC GAGAATGAAT TGCCCCAGGA GTGGGAGAAC AACAAAGACG	780
40	CCCTCCTCAC CTACCTGGAG CAGGTGCGCA TGGGCATTGC AGGAGTGGTG AGGGACAAGG	840
	ACACGGAGCT TGGGATTGCT GACGCTGTCA TTGCCGTGGA TGGGATTAAC CATGACGTGA	900
45	CCACGGCGTG GGGCGGGGAT TATTGGCGTC TGCTGACCCC AGGGGACTAC ATGGTGACTG	960
	CCAGTKCCGA GGGCTACCAT TCAGTGACAC GGAAGTGTGC GGTACCTTT GAAGAGGGCC	1020
	CCTTCCCCTG CAATTTCTG CTCACCAAGA CTCCCAAACA GAGGCTGCGC GAGCTGCTGG	1080
50	CAGCTGGGGC CAAGGTGCCC CCGGACCTTC GCAGGCGCCT GGAGCGGCTA AGGGGACAGA	1140
	AGGATGATA CCTGCGGTTT AAGAGCCCTA GGGCAGGCTG GACCTGTCAA GACGGGAAGG	1200
55	GGAAGACTAG AGAGGGAGGG ACAAAGTGAG GAAAAGGTGC TCATTAAAGC TACCGGGCAC	1260
	CTTAAAAAAA AAAAAAAAAA AAAAA	1285

(2) INFORMATION FOR SEQ ID NO: 85:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 394 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 85:

10 CGCGCTCTA GGAAGTAGTG GATCCCCCGG GNCTGCAGGT GTGGAGTGGG CCATCGTAAA 60
 TAGTATCTGT GCATAAGGTG GTTGTGCGAT AAATGAGTTA ATGTATGCAA AGCCCTTGGC 120
 15 CCAGAGCCGG CGCAGAGCAT TGTGTAAGTS CTGGCAGGCG TCATGATGGA GATATCATGT 180
 CTCCTCTTCT TGATTCAGGA TTCTGATGAG ATGGAGGATG GGCCTGGGGT TCAGGATTAG 240
 GCCTTGAGGC ACTGCTCCAG CCTCCTTTGT GGGCCCTGTC ACCCTTGCT TCATCGGGCC 300
 20 GTARCAAGTC TCCCCTCTCC CACTYTGAG CAGARGTGT CAAGAACTGC CTGCTCACGG 360
 TTCGTGTCT GCAAGGCCAT CGCCTAACCT CTAA 394
 25

(2) INFORMATION FOR SEQ ID NO: 86:

(i) SEQUENCE CHARACTERISTICS:

- 30 (A) LENGTH: 1925 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 35 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 86:

AGTGAAGGGA GCTGGCCGTG CGACTGGGCT TCGGGCCCTG TGCCAGAGGA GCANGCCTTC 60
 40 CTGAGCAGGA GGAAGCAGGT GGTGGCCGCG GCCTTGAGGC AGGCCCTGCA GCTGGATGGA 120
 GACCTGCAGG AGGATGAGAT CCCAGTGGTA GCTATTATGG CCACTGCTGG TGGGATCCGG 180
 GCAATGACTT CCCTGTATGG GCAGCTGGCT GGCCTGAAGG AGCTGGGCCT CTTGGATTGC 240
 45 KTCTCCTACA TCACCGGGGCT CCGGGCTCC ACCTGGGCCT TGGCCAACCT TTATAAGGAC 300
 CCAGAGTGGT CTCAGAAGGA CCTGGCAGGG CCCACTGAGT TGCTGAAGAC CCAGGTGACC 360
 50 AAGAACAAGC TGGGTGTGCT GGCCCCCAGC CAGCTGCAGC GGTACCGGCA GGAGCTGGCC 420
 GAGCGTGCCC GCTTGGGCTA CCCAAGCTGC TTCACCAACC TGTGGGCCCT CATCAACGAG 480
 GCGCTGCTGC ATGATGAGCC CCATGATCAC AAGCTCTCAG ATCAACGGGA GGGCCTGAGT 540
 55 CATGGCCAGA ACCCTCTGCC CATCTACTGT GCCCTCAACA CCAAGGGGA GAGCCTGACC 600
 ACTTTTGAAT TTGGGGAGTG GTGCGAGTTC TCTCCCTACG AGGTGGGCTT CCCCAGTAC 660
 60 GGGGCCTTCA TCCCCTCTGA GCTCTTTGGC TCCGAGTTCT TTATGGGGA GCTGATGAAG 720

	AGGCTTCCTG AGTCCCGCAT CTGCTTCTTA GAAGGTATCT GGAGCAACCT GTATGCAGCC	780
5	AACCTCCAGG ACAGCTTATA CTGGGCCTCA GAGCCCAGCC AGTTCTGGGA CCGCTGGGTC	840
	AGGAACCAGG CCAACCTGGA CAAGGAGCAG GTCCCCCTTC TGAAGATAGA AGAACCACCC	900
	TCAACAGCCG GCAGAATAGC TGAGTTTTC ACCGATCTTC TGACGTGGCG TCCACTGGCC	960
10	CAGGCCACAC ATAATTTCTT GCGTGGCCTC CATTTCCACA AAGACTACTT TCAGCATCCT	1020
	CACTTCTCCA CATGGAAGC TACCACTCTG GATGGGCTCC CCAACCAGCT GACACCCTCG	1080
15	GAGCCCCACC TGTGCCTGCT GGATGTTGGC TACCTCATCA ATACCAGCTG CCTGCCCCTC	1140
	CTGCAGCCCA CTCGGGACGT GGACCTCATC CTGTCAATGG ACTACAACCT CCACGGAGCC	1200
	TTCCAGCAGT TGCAGCTCCT GGGCCGGTTC TGCCAGGAGC AGGGGATCCC GTTCCCACCC	1260
20	ATCTCGCCCA GCCCCGAAGA GCAGCTCCAG CCTCGGGAGT GCCACACCTT CTCCGACCCC	1320
	ACCTGCCCCG GAGCCCCTGC GGTGCTGCAC TTCTCTCTGG TCAGCGACTC CTTCCGGGAG	1380
25	TACTCGGCCC CTGGGGTCCG GCGGACACCC GAGGAGGCGG CAGCTGGGGA GGTGAACCTG	1440
	TCTTCATCGG ACTCTCCCTA CCACTACACG AAGGTGACCT ACAGCCAGGA GGACGTGGAC	1500
	AAGCTGCTGC ACCTGACACA TTACAATGTC TGCAACAACC AGGAGCAGCT GCTGGAGGCT	1560
30	CTGCGCCAGG CAGTGCAGCG GAGGCGGCAG CGCAGGCCCC ACTGATGGCC GGGGCCCCTG	1620
	CCACCCCTAA CTCTCATTCA TTCCCTGGCT GCTGAGTTGC AGGTGGGAAC TGTCATCACG	1680
35	CAGTGCTTCA GAGCCTCGGG CTCAGGTGGC ACTGTCCCAG GGTCCAGGCT GAGGGCTGGG	1740
	AGCTCCCTTG CGCCTCAGCA GTTTGCAGTG GGGTAAGGAG GCCAAGCCCA TTTGTGTAAT	1800
	CACCCAAAAC CCCCCGGCCT GTGCCTGTTT TCCCTTCTGC GCTACCTTGA GTAGTTGGAG	1860
40	CACTTGATAC ATCACAGACT CATACAAATG TGAGGCGCTG AGAAAAAAAA AAAAAAAAAA	1920
	CTCGA	1925

45

(2) INFORMATION FOR SEQ ID NO: 87:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 1818 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 87:

	CCGGGCCCCC CCNCGNGNTT TTTTTTTTTT TTTTTTTTTH TATGAGTCTG TRATGTATCA	60
60	AGTGCTCCAA CTA CTACTCAAGG TAGCGCAGAA GGGAAAACAG GCACAGGCCG GGGGGTTTTG	120

	GGTGATTACA CAAATGGGCT TGGCCTCCTT ACCCCACTGC AAAGTGTGA GGCGCAAGGG	180
	AGCTCCCAGC CCTCAGCCTG GACCCTGGGA CAGTGCCACC TGAGCCCGAG GCTCTGNAAG	240
5	CACTGCGTGA TGACAGTTCC CACCTGCAAC TCAGCAGCCA GGAATGAAT GAGAGTTAGG	300
	GGTGGCAGGG GCCCCGGCCA TCAGTGGGGC CTGCGCTGCC GCCTCCGCTG CACTGCCTGG	360
10	CGCAGAGCCT CCAGCAGCTG CTCCTGGTGT TTGCAGACAT TGTAATGTGT CAGGTGCAGC	420
	AGCTTGTCCA CGTCCTCCTG GCTGTAGGTC ACCTTCGTGT AGTGGTAGGG AGAGTCCGAT	480
	GAAGACAGGT TCACCTCCCC AGCTGCCGCC TCCTCGGGTG TCCGCCGGAC CCCAGGGGCC	540
15	GAGTACTCCC GGAAGGAGTC GCTGACCAGA GGAAAGTGCA GCACCGCAGG GGCTCCGGGG	600
	CAGGTGGGGT CGGAGAAGGT GTGGCACTCC CGAGGCTGGA GCTGCTCTTC GGGGCTGGGC	660
20	GAGATGGGTG GGAACGGGAT CCCCTGCTCC TGGCAGAACC GGCCCAGGAG CTGCAACTGC	720
	TGGAAGGCTC CGTGGAGGTT GTAGTCCAAT GACAGGATGA GGTCCACGTC CCGAGTGGGC	780
	TGCAGGAGGG GCAGGCAGCT GGTATTGATG AGGTAGCCAA CATCCAGCAG GCACAGGTGG	840
25	GGCTCCGAGG GTGTCAGCTG GTTGGGGAGC CCATCCAGAG TGGTAGCTTT CCATGTGGAG	900
	AAGTGAGGAT GCTGAAAGTA GTCTTTGTGG AAATGGAGGC CACGCAGGAA ATTATGTGTG	960
30	GCCTGGGCCA GTGGACGCCA CGTCAGAAGA TCGGTGAAAA ACTCAGCTAT TCTGCCGGCT	1020
	GTTGAGGGTG GTTCTTCTAT CTTCAGAAGG GGGACCTGCT CTTGTCCAG GTTGGCCTGG	1080
	TTCTTGACCC AGCGGTCCCA GAACTGGCTG GGCTCTGAGG CCCAGTATAA GCTGTCTCTG	1140
35	AGGTTGGCTG CATAAGGTT GCTCCAGATA CCTTCTAAGA AGCAGATGCG GGAATCAGGA	1200
	AGCCTCTTCA TCAGCTGCCC CATAAAGAAC TCGAGGCCAA AGAGCTCAGA GGGGATGAAG	1260
40	GCCCCGTAAT TGGGGAAGCC GACCTCGTAG GGAGAGAACT CGCACCCTC CCCAAATTCA	1320
	AAAGTGGTCA GGCTCTGCCC TTTGGTGTG AGGGCACAGT AGATGGGCAG AGGGTTCTGG	1380
	CCATGACTCA GGGCCTCCCG TTGATCTGAG AGCTTGTGAT CATGGGGCTC ATCATGCAGC	1440
45	AGCGCCTCGT TGATGAGGGC CCACAGGTTG GTGAAGCAGC TTGGGTAGCC CAAGCGGGCA	1500
	CGCTCGGCCA GCTCCTGCCG GTACCGCTGC AGCTGGCTGG GGGCCAGCAC ACCCAGCTTG	1560
50	TTCTTGGTCA CCTGGGTCTT CAGCAACTCA GTGGGCCCTG CCAGGTCCTT CTGAGACCAC	1620
	TCTGGGTCTT YATAAAGGTT GGCCAAGGCC CAGGTGGAGC CCGAGGCCCC GGTGATGTAG	1680
	GAGACGCAAT CCAAGAGGCC CCAGCTCCTT TCAGGCCAGC CAGCTGCCCA TACAGGGAAG	1740
55	TCATTGCCCC GATCCACCA CCAGTGGCCA TAATAGCTAC CACTGGGATC TCATCCTCCT	1800
	GCAGGTCTCC ATCCAGCT	1818

(2) INFORMATION FOR SEQ ID NO: 88:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 539 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 88:

10 AGGGTAATTA ATATGAAGTG CAAAAAGTTG AATGTTCCAG TCTAAAAGGC AGTGGGAGAA 60
 ATTACATAGC ATGGAAATAA TAAAATGAAY TCTTATTAAT GAGAACGAGG YTCTTGCACT 120
 15 GGCAAGTTCT GCTGGTCACC CGATGGGGAT GGGAGCCTTT CAAGCTTTTT TTTGGGTAAT 180
 ACTCACAGTT TCCAACGTCT GTGTACTTTT CAAAATGAGC TTGTTCTTCC TTCTGACACT 240
 20 CATCTCAAAG CTCCATGGTG ACGCAGAGGT CTGTTGAAGG TCACAGGGTC CTCGCTTGCA 300
 TTGGCATAAG GTCCTGTAGC ATCACTTGTT AGCCCACTGC TGCTTGAAGG AACTAAGAGT 360
 ATTCAGGGAT AGAGAGCTGA AAATAGGATT AATTNNTTCC TTTTGACTCT CCCCTCAAGA 420
 25 TGTCTTGCT TTTGTTCTGAA AACCTCTCCT GACAACTTTT GCCCAAAGCA AACCATCTGC 480
 CTTTTCTGAA CTCTGAGTGA ATATATTAGC ATCTTCCTT CTGAGCCCTC GTACTGCCA 539
 30

(2) INFORMATION FOR SEQ ID NO: 89:

(i) SEQUENCE CHARACTERISTICS:

- 35 (A) LENGTH: 855 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 89:

40 CCTCTGCCCA GGCCGCACCC GAGCTCAGGC TCGTGCCAC CCACCAAGTT CCAGTGCCGC 60
 45 ACCAGTGGCT TATGCGTGCC CCTCACCTGG CGCTGCGACA GGNACTTGGA CTGCAGCGAT 120
 GGCAGCGATG AGGAGGAGTG CAGGATTGAG CCATGTACCC AGAAAGGGCA ATGCCCCCGG 180
 50 CCCCCTGGCC TCCCCTGCCC CTGCACCGGC GTCAGTGA CTCTGCGGGG AACTGACAAG 240
 AACTGCGCA ACTGCAGCCG CCTGGCCTGC CTAGCAGCGG AGCTCCGTG CACGCTGAGC 300
 GATGACTGCA TTCCACTCAC GTGGCGCTGC GACGGCCACC CAGACTGTCC CGACTCCAGC 360
 55 GACGAGCTCG GCTGTGGAAC CAATGAGATC CTCCCGAAG CGGATGCCAC AACCATGGGG 420
 CCCCCTGTGA CCCTGGAGAG TGTCACCTCT CTCAGGAATG CCACAACCAT GGGGCCCCCT 480
 GTGAACCCTG GAGAGTGTCC CCTCTGTCCG GAATGCCACA TCCCTGCTCG CCGGAGACCA 540
 60

233

GTCTGGAAGC CCAACTGCCT ATGGGGTTAT TGCAGCTGCT GCGGTGCTCA GTGCAAGCCT 600
GGTCACCGCC ACCCTCCTCC TTTTGTCTTG GCTCCGAGCC CAGGAGCGCC TCCGCCCACT 660
5 GGGGTTACTG GTGGCCATGA AGGAGTCCCT GCTGCTGTCA GAACAGAAGA CCTCGCTGCC 720
CTGAGGACAA GCACTTGCCA CCACCGTCAC TCAGCCCTGG GCGTACNGSA CAGGAGGAGA 780
GCAGTGATGC GGATGGGTAC CGGGCACACC AGCCCTTCAG AGACCTGAGC NCTTCTGGCC 840
10 ACTGGAACCTT CGAAC 855

15

(2) INFORMATION FOR SEQ ID NO: 90:

(i) SEQUENCE CHARACTERISTICS:
20 (A) LENGTH: 628 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 90:

AAGGACGTGC CGTGCCGCTG GGTTCCTGAGC CGGAGTGGTC GGTGGGTGGG ATGGAGGCGA 60
CCTTGAGCA GCACTTGAA GACACAATGA AGAATCCCTC CATTGTTGGA GTCCTGTGCA 120
30 CAGATTCACA AGGACTTAAT CTGGGTGCGC GCGGGACCTC GTCAGATGAG CATGCTGGAG 180
TGATATCTGT TCTAGCCCAG CAAGCAGCTA AGCTAACCTC TGACCCCACT GATATTCCTG 240
TGGTGTGTCT AGAATCAGAT AATGGGAACA TTATGATCCA GAAACACGAT GGCATCACGG 300
35 TGGCAGTGCA CAAAATGGCC TCTTGATGCT CATATCTGTT CTTGAGCAGC CTGTCATAGG 360
AACTGGATCC TACCTATGTT AATTACCTTA TAGAACTACT AAAGTTCCAG TAGTTAGGCC 420
40 ATTCATTTAA TGTGCATTAG GCACTTTTCT GTTTATTTAA GAGTCAATTG CTTTCTAATG 480
CTCTATGGAC CGACTATCAA GATATTAGTA AGAAAGGATC ATGTTTTGAA GCAGCAGGTC 540
CAGGTCACCT TGTATATAGA ATTTGCTGT ATTCAATAAA TCTGTTTGGA GGNAAAAAAA 600
45 AAAAAAARAAA AAMTSGAGGG CCGAAGCT 628

50

(2) INFORMATION FOR SEQ ID NO: 91:

(i) SEQUENCE CHARACTERISTICS:
55 (A) LENGTH: 1053 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

60

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 91:

	CTCTTTTCTG CAGTTCAAGG GAAGG ATCTTGCACA AGGCACTCTG CTTCTGCCCT	60
	TGGCTGGGGA AGGGTGGCAT GGAGCTCTC CGGCTGCTCA TCTTACTCTT TGTACAGAG	120
5	CTGTCCGGAG CCCACAACAC CACAGTGTTC CAGGGCGTGG CGGGCCAGTC CCTGCAGGTG	180
	TCTTGGCCCT ATGACTCCAT GAAGCACTGG GGGAGGCGCA AGGCCTGGTG CCGCCAGCTG	240
10	GGAGAGAAGG GCCCATGCCA GCGTGTGGTC AGCACGCACA ACTTGTGGCT GCTGTCCTTC	300
	CTGAGGAGGT GGAATGGGAG CACAGCCATC ACAGACCATA CCCTGGGTGG CACTCTCACC	360
	ATTACGCTGC GGAATCTACA ACCCCATGAT GCGGGTCTCT ACCAGTGCCA GAGCCTCCAT	420
15	GGCAGTGAGG CTGACACCCT CAGGAAGGTC CTGGTGGAGG TGCTGGCAGA CCCCCTGGAT	480
	CACCGGGATG CTGGAGATCT CTGGTTCCCC GGGGAGTCTG AGAGCTTCGA GGATGCCCAT	540
20	GTGGAGCACA GCATCTCCAG GAGCCTCTTG GAAGGAGAAA TCCCCTTCCC ACCCACTTCC	600
	ATCCTTCTCC TCCTGGCCTG CATCTTTCTC ATCAAGATTC TAGCAGCCAG CGNCCTCTGG	660
	GCTGCAGCCT GGCATGGACA GAAGCCAGGG ACACATCCAC CCAGTGAACCT GGACTGTGGC	720
25	CATGACCCAG GGTATCAGCT CCAAACCTCTG CCAGGGCTGA GAGACACGTG AAGGAAGATG	780
	ATGGGAGGAA AAGCCCAGGA GAAGTCCCAC CAGGGACCAG CCCAGCCTGC ATACTTGCCA	840
30	CTTGGCCACC AGGACTCCTT GTTCTGCTCT GGCAAGAGAC TACTCTGCCT GAACACTGCT	900
	TCTCCTGGAC CCTGGAAGCA GGGACTGGTT GAGGGAGTGG GGAGGTGGTA AGAACACCTG	960
	ACAACTTCTG AATATTGGAC ATTTTAAACA CTTACAAATA AATCCAAGAC TGTCATATTT	1020
35	AAAAAAAAA AAAAAAAAAA AACNCGAGGG GGC	1053

40 (2) INFORMATION FOR SEQ ID NO: 92:

(i) SEQUENCE CHARACTERISTICS:

- 45 (A) LENGTH: 1075 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 92:

50	GCACGAGCCT GATCCTCTCT TTTCTGCAGT TCAAGGGAAA GACGAGATCT TGCACAAGGC	60
	ACTCTGCTTC TGCCCTTGGC TGGGGAAGGG TGGCATGGAG CCTCTCCGGC TGCTCATCTT	120
55	ACTCTTTGTC ACAGAGCTGT CCGGAGCCCA CAACACCACA GTTCTTCCAG CAGTGGCGGG	180
	CCAGTCCCTG CAGGTGTCTT GCCCCTATGA CTCCATGAAG GACTTGGGGA GCGGCAAGGC	240
	CTGGTGCCGC CAGCTGGGAG AGAAGGGCCC ATGCCAGCGT GTTCTTCCAG CACACAACCT	300
60	GTGGCTGCTG TCCTTCCTGA GGAGGTGGAA TGGGAGCACA GGCATCACAG ACGATACCCT	360

GGGTGGCACT CTCACCATTA CGCTGCGGAA TCTACAACCC CATGATGCGG GTCTCTACCA 420
 5 GTGCCAGAGC CTCCATGGCA GTGAGGCTGA CACCCTCAGG AAGGTCTCTG TGGAGGTGCT 480
 GGCAGACCCC CTGGATCACC GGGATGCTGG AGATCTCTGG TCCCCCGGG AGTCTGAGAG 540
 CTTTCGAGGAT GCCCATGTGG AGCACAGCAT CTCCAGGAGC CTCTTGGAAG GAGAAATCCC 600
 10 CTTCCACCCC ACTTCCATCC TTCTCCTCCT GGCCTGCATC TTTCTCATCA AGATTCTAGC 660
 AGCCAGCGCC CTCTGGGCTG CAGCCTGGCA TGGACAGAAG CCAGGGACAC ATCCACCCAG 720
 15 TGAAC TGGAC TGTGGCCATG ACCCAGGGTA TCAGCTCCAA ACTCTGCCAG GGCTGAGAGA 780
 CACGTGAAGG AAGATGATGG GAGGAAAAGC CCAGGAGAAG TCCCACCAGG GACCAGCCCA 840
 GCCTGCATAC TTGCCACTTG GCCACCAGGA CTCCTTGTTT TGCTCTGGCA AGAGACTACT 900
 20 CTGCCTGAAC ACTGCTTCTC CTGGACCCTG GAAGCAGGGA CTGGTTGAGG GAGTGGGGAG 960
 GTGGTAAGAA CACCTGACAA CTTCTGAATA TTGGACATTT TAAACACTTA CAAATAAATC 1020
 25 CAAGACTGTC ATATTTAAAA AAAAAAAAAA AAAAAAACN CGAGGGGGGN CCCGG 1075

30 (2) INFORMATION FOR SEQ ID NO: 93:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2492 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:

40 TCCCGACTCA GCTTCCCACC CTGGGCTTTC CGAGGTGCTK TCGCCGCTGT CCCCACTACT 60
 GCAGCCATGA TCTCCTTAAC GGACACGCAG AAAATTGGAA TGGGATTAAC AGGATTGGA 120
 GTGTTTTTCC TGTCTTTGG AATGATCTC TTTTTTGACA AAGCACTACT GCCTATTGGA 180
 45 AATGTTTTAT TTGTAGCCGG CTGGCTTTT GTAATTGGTT TAGAAAGAAC ATTCAGATTC 240
 TTCTTCCAAA AACATAAAAT GAAAGCTACA GGTTTTTTTC TGAATGCTGT ATTTGTAGTC 300
 50 CTTATTGGTT GGCCTTTGAT AGGCATGATC TTCGAAATTT ATGATTTTTT TCTCTGTTC 360
 AGGGGCTTCT TTCCTGTCGT TGTGGCTTT ATTAGAAGAG TGGATCCCTC 420
 CTAAATTTAC CTGGAATTAG ATCATTGTGA GATAAAGTTG GAATAATTA CAATATGGTA 480
 55 TAACAACAAG TGAATTGAA GACTCATTGA AAATATTCTG TTTTATATAA ATTGATTTGA 540
 AGAATATTCA GCACAAAATT AAATTACATG AAATAGCTTG TTTTATTTT TACAGGAGTT 600
 60 TAAACGTAT AGCCTACAAA GTACCAGCAG CAAATTAGCA AAATAATTT GAAAACAGGC 660

	TTCTACTCAA GTGAACTAAG AAGAAGTCAG CAAGCAAACCT GAGAGAGGTG AAATCCATGT	720
	TAATGATGCT TAAGAAACTC TTGAAGGCTA TTTGTGTGTG TTTTCCACAA TGTGCGAAAC	780
5	TCAGCCATCC TTAGAGAACT GTGGTGCCTG TTTCTTTTCT TTTTATTTTG AAGGCTCAGG	840
	AGCATCCATA GGCATTTGCT TTTTAGAAAT GTCCACTGCA ATGGCAAAAA TATTTCCAGT	900
10	TGCACTGTAT CTCTGGAAGT GATGCATGAA TTCGATTGGA TTGTGTCATT TTAAAGTATT	960
	AAAACCAAGG AAACCCCAAT TTTGATGTAT GGATTACTTT TTTTGTATAA CATGGTTAAA	1020
	ATAAACTTC TGTGGTTCTT CTGAATCTTA ATATTTCAAA GCCAGGTGAA AATCTGAACT	1080
15	AGATATTCTT TGTGGAATA TGCAAAGGTC ATTCTTTACT AACTTTTAGT TACTAAATTA	1140
	TAGCTAAGTT TTGTGAGCAG CATACTCCGG AAAGTCTCAT ACTTCTTGGG AGTCTGCCCT	1200
20	CCTAAGTATC TGTCTATATC ATTCATTACG TGTAAGTATT TAACAAAAAA GCATTCTTGA	1260
	CCATGAATGA AGTAGTTTGT TTCATAGCTT GTCTCATTTGA ATAGTATTAT TGAAGATACT	1320
	AAATGATGCA AACCAAATGG ATTTTTCCTA TGTGATGATG TAATTTTCTT TCTTCTTTC	1380
25	TTTTTTTTAA ATTTTAGCAG TGGCTTATTA TTTGTTTTTC ATAAATTAAA ATAACTTTTG	1440
	ATAATGTTTA CTTTAAGACA TGTAACATGT TAAAAGGTTA AACTTATGGC TGTTTTTAAA	1500
30	GGGCTATTCA TTTAATCTGA GTTTTCCCTT ATTTTCAGCT TTTTCCTAGC ATATAATAGT	1560
	CATTAAGCAT GACATATCCT TCATATGATC ACTCATCTTG AGTTAATTAG AAAATACCTG	1620
	AGTTCACGTG CTAAAGTCAT TTCACTGTAA TAACTGACT RTGGTTTCTT AAGAACATGA	1680
35	CACTAAAAAA AAAGTGGTTT TTTTCCACCG TTGCTGATTA TTAGACAGTA GGAAATAGCT	1740
	GTTTCTTTTA GTTTTACAAG ATGTGACAGC TTTAGTGGTA GATGTAGGGA AACATTTCAA	1800
40	CAGCCATAGT ACTATTGTGTT TTACCACTGA TTGCACTGTT TTGTTTTTTT AACAGTTGCA	1860
	AAGCTTTTTA ATGCATAAAA GTATAATTGA AATCTGTGGT ATTTATTTAC AAACATGTCT	1920
	ACAAAAATAG ATTACAGCTT ATTTTATTTT TAGTTAAATC TCTTAATACA CAGAGNAACT	1980
45	CCCAATCTTG CTCATCTAAA TAAGGAAAGA CTTGGTGTAT AGTGTGATGG TTTAGTCTTA	2040
	AGGATTAAGA CATTTTTGGT ACTTGCATTT GACTTACGAT GTATCTGTGA AAATGGGATG	2100
50	ATATTGACAA ATGGAGACTC CTACCTCAAT AGTTAATGGA ATATAAGAG GCTACTGTTG	2160
	TGTCTAATGT TCTTCAAAAA AGTAATATCC TCACTTGGAG AGTGTGAAAT ACATACTTTG	2220
	AGGATTGACT TTATATAAGG TGCCCTGTAG AAMTCTGTTA TATATATCTT TGACCCATAT	2280
55	TATTTACAAT GTCTTGATAA TTCTACCTTT TTAGAGCAAG AATAATGTTT TTAAATGTAA	2340
	GGGACATCTG TATTTAACTC CTTTGTAGAC ATGAATTTCT ATTAATAATC TCTTTGCACT	2400
60	GTAACAGAGA TTCCTTTTTT CAATAATCTT AATTCAAAGC ATATTAATGTTT GTTGAAAGGG	2460

TTTGRTAATC TCCCCGTCCT TGGTAAAGGT TG

2492

5

(2) INFORMATION FOR SEQ ID NO: 94:

(i) SEQUENCE CHARACTERISTICS:

10

- (A) LENGTH: 3058 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 94:

15

ACCCATAATC AACAGACAAT GGCATTGTCTG AAGAGCAACC TGTTAATGAA ATCATGTTAA 60

AAATCAAGGT TTGGCTTCAG TTAAATCAC TTGAGGTATG AAGTTTATCC TGTTTTCCAG 120

20

AGATAAACAT AAGTTGATCT TCCAAAATA CCATCATTAG GACCTATCAC ACAATATCAC 180

TAGTTTTTTT TGTGTGTTT TTTTTGTTC TTTTCTTGG TAAAGCCATG CACCACAGAC 240

25

TTCTGGGCAG AGCTGAGAGA CAATGGTCCT GACATAATAA GGATCTTTGA TTAACCCCCA 300

TAAGGCATGT GTGTGTATAC AAATATACTT CTCTTTGGCT TTTGACATA GAACCTCAGC 360

TGTTAACCAA GGGGAAATAC ATCAGATCTG CAACACAGAA ATGCTCTGCC TGAAATTTCC 420

30

ACCATGCCTA GGAATCACCC CATTTATCCA GGTCTTTCTG GATCTGTTTA ATCAATAAGC 480

CCTATAATCA CTGCTAAAC ACTGGGCTTC ATCACCAGG GATAAAAACA GAGATCATTG 540

35

TCTTGGACCT CCTGCATCAG CCTATTCAAA ATTATCTCTC TCTCTAGCTT TCCACAAATC 600

CTAAATTTCC TGTCCCAAGC CACCCAAATT CTCAGATCTT TTCTGGAACA AGGCAGAATA 660

TAAAATAAAT ATACATTTAG TGGCTTGGGC TATGGTCTCC AAAGATCCTT CAAAAATACA 720

40

TCAAGCCAGC TTCATTCACT CACTTTACTT AGAACAGAGA TATAAGGGCC TGGGATGCAT 780

TTATTTTATC AATACCAATT TTTGTGGCCA TGGCAGACAT TGCTAATCAA TCACAGCACT 840

45

ATTTCTTAT T AAGCCCACTG ATTTCTTCAC AATCCTTCTC AAATTACAAT TCCAAAGAGC 900

CGCCACTCAA CAGTCAGATG AACCCAACAG TCAGATGAGA GAAATGAACC CTACTTGCTA 960

TCTCTATCTT AGAAAGCAAA AACAAACAGG AGTTTCCAGG GAGAATGGGA AAGCCAGGGG 1020

50

GCATAAAAGG TACAGTCAGG GGAAATAGA TCTAGGCAGA GTGCCTTAGT CAGGGACCAC 1080

GGGCGCTGAA TCTGCAGTGC CAACACCAAA CTGACACATC TCCAGGTGTA CCTCCAACCC 1140

55

TAGCCTTCTC CCACAGCTGC CTACAACAGA GTCTCCAGC CTTCTCAGAG AGCTAAAACC 1200

AGAAATTTCC AGACTCATGA AAGCAACCCC CCAGCCTCTC CCCAAGCCTG CCGCATTTGTC 1260

TAATTTTTAG AACACTAGGC TTCTTCTTTC ATGTAGTTCC TTATAAGCAG GGGCCAGAAT 1320

60

ATCTCAGCCA CCTGCAGTGA CATTGCTGGA CCCCTGAAAA CCATTCCATA GGAGAATGGG 1380

	TTCCCCAGGC TCACAGTGTA GAGACATTGA GCCCATCACA ACTGTTTTGA CTGCTGGCAG	1440
5	TCTAAACAG TCCACCCACC CCATGGCACT GCCGCGTGAT TCCCGCGCCA TTCAGAAGTT	1500
	CAAGCCGAGA TGCTGACGTT GCTGAGCAAS AGATGGTGAG CATCAGTGCA AATGCACCAT	1560
	TCAGCACATC AGTCATATGC CCAGTGCACT TACAAGATGT TGTTCGGCA AAGCATTTTG	1620
10	ATGGAATAGG GAACTGCAAA TGTATGATGA TTTTGAAAAG GCTCAGCAGG ATTTGTTCTT	1680
	AAACCGACTC AGTGTGTCAT CCCCGGTTAT TTAGAATTAC AGTTAAGAAG GAGAAACTTC	1740
15	TATAAGACTG TATGAACAAG GTGATATCTT CATAGTGGGC TATTACAGGC AGGAAAATGT	1800
	TTTAACTGGT TTACAAAATC CATCAATACT TGTGTCATTC CCTGTAAAAG GCAGGAGACA	1860
	TGTGATTATG ATCAGGAAAC TGCACAAAAT TATTGTTTTC AGCCCCCGTG TTATTGTCCT	1920
20	TTTGAACTGT TTTTTTTTTT TAAAGCCAA ATTTGTGTTG TATATATTCG TATTCCATGT	1980
	GTTAGATGGA AGCATTTTCT ATCCAGTGTG AATAAAAAGA ACAGTTGTAG TAAATTATTA	2040
25	TAAAGCCGAT GATATTTTCAT GGCAGGTTAT TCTACCAAGC TGTGCTTGTT GGTTTTTTCCC	2100
	ATGACTGTAT TGCTTTTATA AATGTACAAA TAGTTACTGA AATGACGAGA CCCTTGTTTG	2160
	CACAGCATTA ATAAGAACCT TGATAAGAAC CATATTCTGT TGACAGCCAG CTCACAGTTT	2220
30	CTTGCTGAA GCTTGGTGCA CCCTCCAGTG AGACACAAGA TCTCTCTTTT ACCAAAGTTG	2280
	AGAACAGAGC TGGTGGATTA ATTAATAGTC TTCGATATCT GGCCATGGGT AACCTCATTG	2340
35	TAATATCAT CAGAATGGGC AGAGATGATC TTGAAGTGTC ACATACACTA AAGTCCAAAC	2400
	ACTATGTCAG ATGGGGGTAA AATCCATTAA AGAACAGGAA AAAATAATTA TAAGATGATA	2460
	AGCAAATGTT TCAGCCCAAT GTCAACCCAG TTAATAAATAA AATTAATGCT GTGTAAAATG	2520
40	GTTGAATTAG TTTGCAAACT ATATAAAGAC ATATGCAGTA AAAAGTCTGT TAATGCACAT	2580
	CCTGTGGGAA TGGAGTGTTT TAACCAATTG CCTTTTCTTG TTATCTGAGC TCTCCTATAT	2640
45	TATCATACTC AGATAACCAA ATTAAAAGAA TTAGAATATG ATTTTAAATA CACTTAACAT	2700
	TAAACTCTTC TAACTTTCTT CTTTCTGTGA TAATTCAGAA GATAGTTATG GATCTTCAAT	2760
	GCCTCTGAGT CATTGTTATA AAAAATCAGT TATCACTATA CCATGCTATA GGAGACTGGG	2820
50	CAAAACCTGT ACAATGACAA CCCTGGAAGT TGCTTTTTTTT AAAAAATAA TAAATTTCTT	2880
	AAATCAACTC TTTTTTCTGG TTGTCTGTTT GTTATAAAGT GCAACGKATT CAAGTCCTCA	2940
55	ATATCCTGAT CATAATACCA TGCTATAGGA GACTGGGCAA AACTGTGACA ATGACAACCC	3000
	TGGAAGTTGC TTTTTTAAAA AAATAATAAT TTNPTAATCC AAAAAANAA AAAAANTT	3058

(2) INFORMATION FOR SEQ ID NO: 95:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 1099 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 95:

10 GGCTTTGTAG CTGCTCCGCA GCCCAGCCCG GCGCGCTCG CAGAGTCCTA GGCGGTGCGC 60
 GGCNTCCTGC CTCCTCCCTC CTCGGCGGTC GCGGCCCGCG CCTCCGCGGT GCCTGCCTTC 120
 15 GCTCTCAGGT TGAGGAGCTC AAGCTTGGA AAATGGTGTG CATTCCTTGT ATCGTCATTC 180
 CAGTTCTGCT CTGGATCTAC AAAAAATTCC TGGAGCCATA TATATACCCT CTGGTTTCCC 240
 CCTTCGTTAG TCGTATATGG CCTAAGAAAG CAATACAAGA ATCCAATGAT ACAAACAAAG 300
 20 GCAAAGTAAA CTTTAAGGGT GCAGACATGA ATGGATTACC AACAAAAGGA CCAACAGAAA 360
 TCTGTGATAA AAAGAAAGAC TAAAGAAATT TTCCTAAAGG ACCCCATCAT TTAAAAATG 420
 25 GACCTGATAA TATGAAGCAT CTTCCTTGTA ATTGTCTCTG ACCTTTTAT CTGAGACCGG 480
 AATTCAGGAT AGGAGTCTAG ATATTTACCT GATACTAATC AGGAAATATA TGATATCCGT 540
 ATTTAAATG TAGTTAGTTA TATTTAATGA CCTCATTCCT AAGTTCCTTT TTCGTTAATG 600
 30 TAGCTTTCAT TTCTGTTATT GCTGTTTGAA TAATATGATT AAATAGAAGG TTTGTGCCAG 660
 TAGACATTAT GTTACTAAAT CAGCACTTTA AAATCTTTGG TTCTCTAATT CATATGAATT 720
 35 TGCTGTTTGC TCTAATTTCT TTGGGCTCTT CTAATTTGAG TGGAGTACAA TTTTGTGTG 780
 AAACAGTCCA GTGAACTGT GCAGGGAAAT GAAGGTAGAA TTTTGGGAGG TAATAATGAT 840
 GTGAAACATA AAGATTTAAT AATTACTGTC CAACACAGTG GAGCAGCTTG TCCACAAATA 900
 40 TAGTAATTAC TATTTATTCG TCTAAGGAAG ATTAACAAAAA GATAGGGAAA AGGGGGAAAC 960
 TTCTTTGAAA AATGAAACAT CTGTTACATT AATGTCTAAT TATAAAATTT TAATCCTTAC 1020
 45 TGCATTTCTT CTGTTCTTAC AAATGTATTA AACATTCAGT TTAAGTGGTA AAAAAAAAAA 1080
 AAAAAAACC GGGGGGGG 1099

50

(2) INFORMATION FOR SEQ ID NO: 96:

(i) SEQUENCE CHARACTERISTICS:

- 55 (A) LENGTH: 1580 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

60

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 96:

	GGCAGAGACT GGAATCTCTC TTCATGAAAA AATGCAGCCC CTTAACTTCA GTTCGACARA	60
5	GTGCAGCTCC TTCTCTCCAC CCACCACAGT GATTCTCCTT ATCCTGCTGT GCTTTGAGGG	120
	CCTGCTCTTC CTCATTTTCA CATCAGTGAT GTTTGGGACC CAGGTGCACT CCATCTGCAC	180
	AGATGAGACG GGAATAGAAC AATTGAAAAA GGAAGAGAGA AGATGGGCTA AAAAAACAAA	240
10	ATGGATGAAC ATGAAAGCCG TTTTGGCCA CCCCTTCTCT CTAGGCTGGG CCAGCCCCTT	300
	TGCCACGCCA GACCAAGGGA AGGCAGACCC GTACCAGTAT GTGGTCTGAA GGACCCCGAC	360
15	CGGCATGGCC ACTCAGACAC AAGTCCACAC CACAGCACTA CCGTCCCATC CGTTCTCATG	420
	AATGTTTAAA TCGAAAAAGC AAAACAACCTA CTCTTAAAC TTTTTTTATG TCTCAAGTAA	480
	AATGGCTGAG CATTGCAGAG ARAAAAAAAA GTCCCCACAT TTTATTTTAT AAAAAACATC	540
20	CTTTCGATTT CTMTTGGTGA CCGAWGCTGC TCTCTTTTCC TTTTAAATC ACTTCTCTGG	600
	CCTCTGGTTT CTCTCTGCTG TCTGTCTGGC ATGACTAATG TAGAGGGCGC TGTCTCGCGC	660
25	TGTGCCCCATT CTAATACTG AGTGAGACAT GACGCTGTGC TGGATGGAAT AGTCTGGACA	720
	CCTGGTGGGG GATGCATGGG AAAGCCAGGA GGGCCCTGAC CTCCCCTGAC CCAGGAGGCA	780
	GTGGCGGGCT CCCCGATGGG ACATAAAACC TCACCGAAGA TGGATGCTTA CCCCTGAGG	840
30	CCTGAGAAGG GCAGGATCAG AAGGACCTT GGCACAGCGA CCTCATCCCC CAAGTGACA	900
	CGGTTTGCCT GCTAACTCGC AAAGCAATTG CCTGCCTTGT ACTTTATGGG CTTGGGGTGT	960
35	GTAGAATGAT TTTGCGGGG AGTGGGAGA AAGATGAAAG AGGTCTTATT TGTATTCTGA	1020
	ATCAGCAATT ATATTCCCTG TGATPATTTG GAAGAGTGTG TAGGAAAGAC GTTTTTCAG	1080
	TTCAAAATGC CTTATACAAT CAAGAGGAAA AAAAATTACA CAATTTACAG CAAGCTACGT	1140
40	TTTCCTTTGT TTCATCTGCT TCCTCTCTCA CCACCCCATC TCCCTCTCTT CCCCAGCAAG	1200
	ATGTCAATTA AGCAGTGTGA ATTCTGACTG CAATAGGCAC CAGTGCCCAA CACATACAGC	1260
45	CCCACCATCA TCCCCTTCTC ATTTTATAAA CCTCAAAGTG GATTCACTTT CTGATAGTTA	1320
	ACCCCCATAA ATGTGCACGT ACCTGTGTCT TATCTATATT TTAACCKGGG AGACTGTTGT	1380
	CCTGGGCATG GGAGATGACC ATGATGCTGG GGTACCTCA CAGTCCCCAC CCTTTCAAAG	1440
50	TTNGACATAT GGGCCATCCC ATTGGGCCAG GAATTCCACA GGACACACCT AAGGCTGTGG	1500
	GMAYTGGGGG ACAAATAGAT TTTCCATTTT GAGGAGGCA CTTTCCCTGT TGTTCACTTC	1560
55	TTGTTTTGAA GGGAGGTNGG	1580

(2) INFORMATION FOR SEQ ID NO: 97:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 678 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 97:

10 ATATTTTTTT AGGCTAATGT CCAAGATACA GCATTGAGGA GGCAGCTATG TCTAATGAGG 60
 GCTCTCTTGT TTGCTAGAGA TGAGAGAAAT GTATACTAAT CATTTTAATT TGTACTTAAA 120
 ATACATTTTA CTAATCATAT TGATTTTAAA TATGACAAAT TCTTCTAGTA GATACTAATC 180
 15 TTTCTTGTTT ATCATATTGT CCTAGAGAAG CCTAGGTAAA AATGGGTTC ACCTAGTCTG 240
 TTTGTATAAC ACCTTCCCC GTCCCCTCTC CATCCCTGCC AATGGGCTC TATGCATATT 300
 20 GACAAGCAAA TAAGAAAACC TTAGGTTTCT TGTATTTGAA TTTCCAAAAC AATAAAAGGT 360
 TTTGACTCAA GATTTGCATT CAAGAAGAGG CAGAAATTTT GTCTTATCTT TTTATCATTT 420
 TGTGAACTTG TGTTTCTCTG TATGCTTAGA AAATTTTACA CACAAGGAAT GTTTGAAAAA 480
 25 GTGAGAATTT TAGAGTGCTT GGGTGGTTTT TATTTGGTCA GTGCTGATGT GTTARGTGTT 540
 TAGGGAAATA ATGCTTCAGG ACCTTTTTGA CAACACAGYT TCATGAATGA CYGGGGGATA 600
 30 TTWAKGTTGT GCTGAGAAAA GGGAGGGAGT GGGCAGTTGG AATGGGGGAC CCTTACCATT 660
 GGAAAACATG CATTCTNGN 678

35

(2) INFORMATION FOR SEQ ID NO: 98:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1253 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

40

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 98:

45

ACCTCCCTCC CTCTCAGACT GGTCGAATC CACGCCTAGC CCAGCCACTG CCACTGGGGC 60
 CATGGCCACC ACCACTGGGG CACTGCCTGC CCAGCCACTT CCCTTGCTCTG TTCCCAGCTC 120
 50 CCTTGCTCAG GCCCAGACCC AGCTGGGGCC CCACCGGNAA GTTACCCCCA AGAGGCAAGT 180
 NTTGGCCTGA GACGCTCGTC AGTTCTTAGA TCTTGGGGGC CTAAAGAGAC CCCCCTCCTG 240
 CCTCCTTTCT TTCTCTGTCT CTTCTTCCT TTTAGTCTTT TTCATCTCTT TCTCTTTCCA 300
 55 CCAACCCTCC TGCATCCTTG CCTTGCAGCG TGACCGAGAT AGGTGATCAG CCCAGGGCTT 360
 CAGTCTTCCT TTATTTATAA TGGGTGGGGG CTACCACCCA CCGTCTCTGA CTCTTGTAAG 420
 60 GAGTCTGGGA CCTCCTTCTT CCCCCTTCT CTCTTCCCTC ATTCTTTCTT CTCTCCTTCT 480

5 GGCCTCTCAT TTCCTTACAC TCTGACATGA ATGAATTATT ATTATTTTTC TTTTCTTTT 540
 TTTTTTTACA TTTTGTATAG AAACAAATTC ATTTAAACAA ACTTATTATT ATTATTTTTC 600
 ACAAATATA TATATGGAGA TGCTCCCTCC CCCTGTGAAC CCCCAGTGC CCCCCTGGGC 660
 TGNAGTCTGT GGGCCCATTC GGCCAAGCTG GATTCGTGT ACCTAGTACA CAGGCATGAC 720
 10 TGGGATCCCG TGTACCGAGT ACACGACCCA GGTATGTACC AAGTAGGCAC CCTTGGGCGC 780
 ACCCACTGGG GCCAGGGGTC GGGGAGTGT TGGGAGCCTC CTCCCCACCC CACCTCCCTC 840
 ACTTCACTGC ATTCCAGATT GGACATGTTC CATAGCCTTG CTGGGAAGG GCCCACTGCC 900
 15 AACTCCCTCT GCCCCAGCCC CACCCTTGGC CATCTCCCTT TGGGAAGTAG GGGGCTGCTG 960
 GTGGGAAATG GGAGCCAGGG CAGATGTATG CATTCCTTTA TGTCCCTGTA AATGTGGGAC 1020
 20 TACAAGAAGA GGAGCTGCCT GAGTGGTACT TTCTCTTCCT GGTAATCCTC TGGCCCAGCC 1080
 TTATGGCAGA ATAGAGGTAT TTTTAGGCTA TTTTGTAAAT ATGGCTTCTG GTCAAAATCC 1140
 CTGTGTAGCT GAATTCCTAA GCCCTGCATT GTACAGCCCC CCACTCCCCT CACCACCTAA 1200
 25 TAAAGGAATA GTTAACACTC AAAAAAAAAA AAAAAAAAAA ACTTGAGGGG GGG 1253

30

(2) INFORMATION FOR SEQ ID NO: 99:

(i) SEQUENCE CHARACTERISTICS:

35

- (A) LENGTH: 447 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 99:

40

CAAAGAATGA AATTTACCAC TCTCCTCTC TTGGCAGCTG TAGCAGGGGC CCTGGTCTAT 60
 GCTGAAGATG CCTCCTCTGA CTCGACGGGT GCTGATCCTG CCCAGGAAGC TGGGACCTCT 120
 45 AAGCCTAATG AAGAGATCTC AGGTCCAGCA GAACCAGCTT CACCCCCAGA GACAACCACA 180
 ACAGCCCAGG AGAYTTCGGC GGCAGCAGTT CAGGGGACAG CCAAGGTCAC CTCAAGCAGG 240
 CAGGAACTAA ACCCCCTGAA ATCCATACTG GAGAAAAGTA TCTTACTAAC AGAACAAGCC 300
 50 CTTGCAAAAG CAGGAAAAGG AATGCACGGA GGCGTGCCAG GTGGAAAACA ATTCATCGAA 360
 AATGGAAGTG AATTGACACA AAAATTACTG AAGAAATCA GTCTATTAAA ACCATGGGCA 420
 55 TGAGAAGCTG AAAAGAATKG GATCATT 447

60

(2) INFORMATION FOR SEQ ID NO: 100:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 611 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 100:

10 GGTCTGGGGA GGTGACATGT TGGGCTGTGG GATCCCAGCG CTGGGCCTGC TCCTGCTGCT 60
GCAGGSWTCG GCAGACGGAA ATGGAATCCA GGGATTCTTC TACCCATGGA GCTGTGAGGG 120
TGACATATGG GACCGGGAGA GCTGTGGGGG CCAGGCGGCC ATTGATAGC CCCAACYTCT 180
15 GCCTGCGTCT CCGGTGCTGC TACCGCAATG GGTCTGCTAC CACCAGCGTC CAGACGAAAA 240
CGTGCGGAGG AAGCACATGT GGGCGCTGGT CTGGACGTGC AGCGGCCTCC TCCTCCTGAG 300
20 CTGCAGCATC TGCTTGTTMT GGTGGGCCAA GCGCCGGGAC GTGCTGCATA TGCCCGGTTT 360
CCTGGCGGGT CCGTGTGACA TGTCCAAGTC CGTCTCGCTG CTCTCCAAGC ACCGAGGGAC 420
CAAGAAGACG CCGTCCACGG GCAGCGTGCC AGTCGCCCTG TCCAAAGAGT CCAGGGATGT 480
25 GGAGGGAGGC ACCGAGGGGG AAGGGACGGA GGAGGGTGAG GAGACAGAGG GCGAGGAAGA 540
GGAGGATTAG GGGAGTCCCC GGGGGACTGG TCAATACAGA TACGGTGGAC GGAAAAAAAA 600
30 AAAAAAAAAA A 611

35 (2) INFORMATION FOR SEQ ID NO: 101:

(i) SEQUENCE CHARACTERISTICS:

- 40 (A) LENGTH: 609 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 101:

45 GCATTGGTAA AGCTGGCAGT TGAAACCAGT TGGACGGCCC AGCTTGCGTC TCTTCTGCCT 60
GAGTGGGCCT CTCAGGTCAC TCGTGCCCTG CTGGAGGACA GAGGGGCACC TCAGCCGCCC 120
CCAAGCCCAG AGCACAGCAA TAAGGTCGGC CTGCAGGAGC CGGGGTGGGG GTGGGGGTGG 180
50 GGGGRGCAGG ACCCTRARAT GCCACCAGGA CCTGATGGGC CAGGAAGGGC GTGGACATGG 240
AGGCTGTTTT TACAGTTTTT TTTTTTTTGT TGTTTTGT TTAAAGAATA CAGAAGGAGC 300
55 CAAGCTTTTT TGCACTTTGT ATCCAGCTGC AAGCTCAGGG CAGAGTCAGG GGGCTGGGTT 360
GGAAAAACCT GACTCACAGG AATGCATAAT TGACCCTTGC AGCTACCCAA TAGCCCTTGG 420
AGCTGGCACT GAACCAGGCT GCAAGATTGT ACTGCCTTAA AAACACAAGG CCCTCTAGGC 480
60

	CTGGCAGGGA TGTCCCTGTG CCCAGCACTG GGGGCTCGAA GACTGGTTTC TAGCACTACC	540
	GGTCACGGCC ATGTCGTCCT AGAAGGGTCC AGAAGATTAT TTTACGTTGA GTCCATTTTT	600
5	AATGTTCTG	609
10	(2) INFORMATION FOR SEQ ID NO: 102:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1770 base pairs	
	(B) TYPE: nucleic acid	
15	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 102:	
20	ACGGYCCGGA ATCCCGGGTC GACCCACGCG TCCGGGAAAT TGAAACTGAG TGGCCCACGA	60
	TGGGAAGAGG GGAAAGCCCA GGGGTACAGG AGGCCTCTGG GTGAAGGCAG AGGCTAACAT	120
	GGGGTTCCGA GCGACCTTGG CCGTTGGCCT GACCATCTTT GTGCTGTCTG TCGTCACTAT	180
25	CATCATCTGC TTCACCTGCT CCGTGTCTG CCTTTACAAG ACGTGCCGCC GACCACGTCC	240
	GGTTGTCACC ACCACCACAT CCACCACTGT GGTGCATGCC CCTTATCCTC AGCCTCCAAG	300
30	TGTGCCGCCC AGCTACCCCTG GACCAAGCTA CCAGGGCTAC CACACCATGC CGCCTCAGCC	360
	AGGGATGCCA GCAGACCCTT ACCCAATGCA GTACCCACCA CCTTACCCAG CCCAGCCCAT	420
	GGGCCCACCG GCCTACCACG AGACCCTGGC TGGAGGAGCA GCCGCGCCCT ACCCCGCCAG	480
35	CCAGCCTCCT TACAACCCGG SCTACATGGA TGCCCCGAAG SGGNCCTCTG AGCATTCCTT	540
	GGCCTCTYTG GCTGCCACTT GGTATGTTG TGTGTGTGCG TGARTGGTGT GCAGGCGCGG	600
40	TTCTTTACGC CCCATGTGTG CTGTGTGTGT CCTGCCTGTA TATGTGGCTT CCTCTGATGC	660
	TGACAAGGTG GGAACAATC CTTGCCAGAG TGGGCTGGGA CCAGACTTTG TTCTCTTCCT	720
	CACCTGAAAT TATGCTTCCT AAAATCTCAA GCCAACTCA AAGAATGGGG TGGTGGGGGG	780
45	CACCTGTGA GGTGGCCCTT GAGAGGTGGG GGCCTCTCCA GGGCACATCT GGAGTTCTTC	840
	TCCAGCTTAC CCTAGGGTGA CCAAGTAGGG CCTGTACAC CAGGGTGGCG CAGCTTTCTG	900
50	TGTGATGCAG ATGTGTCCTG GTTTCGGCAG CGTAGCCAGC TGCTGCTTGA GGCCATGGCT	960
	CGTCCCCGGA GTTGGGGGTA CCCGTGTCAG AGCCAGGGAC ATGATGCAGG CGAAGCTTGG	1020
	GATCTGGCCA AGTTGGACTT TGATCCTTTG GGCAGATGTC CCATTCGCTC CTGGAGCCTG	1080
55	TCATGCCTGT TGGGGATCAG GCAGCCTCCT GATGCCAGAA CAGCTCAGGC AGAGCCCTAC	1140
	TCAGCTGTAC CTGTCTGCCT GGAAGTGTCC CTGTCCCCGC ATCTCCCTTG GGACCAGCTG	1200
60	GAGGGCCACA TGCACACACA GCCTAGCTGC CCCCAGGGAG CTCTGCTGCC CTGCTGGCC	1260

245

CTGCCCTTCC CACAGGTGAG CAGGGCTCCT GTCCACCAGC ACACTCAGTT CTCTTCCCTG 1320
 CAGTGTTTTTC ATTTTATTTT AGCCAAACAT TTTGCCTGTT TTCTGTTTCA AACATGATAG 1380
 5 TTGATATGAG ACTGAAACCC CTGGGTGTG GAGGGAAATT GGCTCAGAGA TGGACAACCT 1440
 GGCAACTGTG AGTCCCTGCT TCCCGACACC AGCCTCATGG AATATGCAAC AACTCCTGTA 1500
 10 CCCAGTCCA CGGTGTTCTG GCAGCAGGGA CACCTGGGCC AATGGGCCAT CTGGACCAA 1560
 GGTGGGGTGT GGGGCCCTGG ATGGCAGCTC TGGCCCAGAC ATGAATACCT CGTGTTCCTC 1620
 CTCCTCTAT TACTGTTTCA CCAGAGCTGT CTTAGCTCAA ATCTGTTGTG TTTCTGAGTC 1680
 15 TAGGGTCTGT ACACTTGTTC ATAATAATG CAATCGTTTG GAAAAAAAAA AAAAAAAAAAC 1740
 TCGTAGGGGG GGCCCGTACC CAATSGCCTA 1770
 20

(2) INFORMATION FOR SEQ ID NO: 103:

25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1832 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear
 30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103:

TGTGGCTGAC GTCATCTGGA GGAGATTTGC TTTCTTTTTC TCCAAAAGGG GAGGAAATTG 60
 35 AAAGTGCAGT GGCCACGAT GGGAAGAGGG GAAAGCCCAG GGGTACAGGA GGCCTCTGGG 120
 TGAAGCAGA GGCTAACATG GGGTTCGGAG CGACCTTGGC CGTTGGCTGA CCATCTTTGT 180
 GCTGTCTGTC GTCACATCA TCATCTGCTT CACCTGCTCC TGCTGCTGCC TTTACAAGAC 240
 40 GTGCCGCCGA CCACGTCCGG TTGTCAACAC CACCACATCC ACCACTGTGG TGCATGCCCC 300
 TTATCCTCAG CCTCCAAGTG TGCCGCCAG CTACCCTGGA CCAAGCTACC AGGGCTACCA 360
 45 CACCATGCCG CCTCAGCCAG GGATGCCAGC AGCACCTAC CCAATGCAGT ACCCACCACC 420
 TTACCCAGCC CAGCCCATGG GCCCACCAGC CTACCACGAG ACCCTGGCTG GAGGAGCAGC 480
 CGCGCCCTAM CCCGSCAGCC AGCCTCCTTA CAACCCGGCC TACATGATG CCCGAAGCGG 540
 50 CCCTCTGAGC ATTCCCTGGC CTCTYTGCT GCCACTTGGT TACTTTTCTG GTGTGCGTRA 600
 GTGGTGTGCA GGCGCGGTTC CTTACGCCCC ATGTGTGCTG TTTTCTTCA AGCAGCGTTC 660
 CTTACGCCCC ATGTGTGCTG TGTGTGCTCT GCCTGTATAT GTTTTCTCT CTGATGCTGA 720
 55 CAAGTGGGGA ACAATCCTTG CCAGAGTGGG CTGGGACCAG ATTTTCTCT CTTCCTCACC 780
 TGAAATTATG CTTCCTAAAA TCTCAAGCCA AACTCAAAGA ATTTTCTCT GGGGGGCACC 840
 60

246

	CTGTGAGGTG GCCCTGAGA GGTGGGGGCC TCTCCAGGGC ACATCTGGAG TTCTTCTCCA	900
	GCTTACCCTA GGGTGACCAA GTAGGGCCTG TCACACCAGG GTGGCGCAST TTCTGTGTGA	960
5	TGCAGATGTG TCCTGGTTTC GGCAGCGTAG CCAGCTGCTG CTTGAGGCCA TGGCTCGTCC	1020
	CCGGAGTTGG GGGTACCCGT TGCAGAGCCA GGGACATGAT GCAGGCGAAG YTTGGGATCT	1080
10	GGCCAAGTTG GACTTTGATC CTTTGGGCAG ATGTCCCAT TCTCCCTGGA GCCTGTCATG	1140
	CCTGTTGGGG ATCAGGCAGC CTCCTGATGC CAGAACACCT CAGGCAGAGC CCTACTCAGC	1200
	TGTACCTGTC TGCCTGGACT GTCCCTGTC CCCGCATCTC CCCTGGGACC AGCTGGAGGG	1260
15	CCACATGCAC ACACAGCCTA GCTGCCCCCA GGGAGCTCTG CTGCCCTTGC TGGCCCTGCC	1320
	CTTCCACAG GTGAGCAGGG CTCCTGTCCA CCAGCACACT CAGTTCTCTT CCCTGCAGTG	1380
20	TTTTCATTTT ATTTTAGCCA AACATTTTGC CTGTTTCTG TTTCAAACAT GATAGTTGAT	1440
	ATGAGACTGA AACCCTGGG TTGTGGAGGG AAATTGGCTC AGAGATGGAC AACCTGGCAA	1500
	CTGTGAGTCC CTGCTTCCCG ACACCAGCCT CATGGAATAT GCAACAAC TCGTACCCCA	1560
25	GTCCACGGTG TTCTGGCAGC AGGGACACCT GGGCCAATGG GCCATCTGGA CCAAAGGTGG	1620
	GGTGTGGGGC CCTGGATGGC AGCTCTGGCC CAGACATGAA TACCTCGTGT TCCTCCTCCC	1680
30	TCTATTACTG TTTCACCAGA GCTGTCTTAG CTCAAATCTG TTGTGTTTCT GAGTCTAGGG	1740
	TCTGTACACT TGTTTATAAT AAATGCAATC GTTTNGGAAA AAAAAANAA AAAAAAAGG	1800
	GGSGGCGCTC TAAAAGGATN CCCNAAGGG GG	1832
35		

(2) INFORMATION FOR SEQ ID NO: 104:

- 40 (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 2237 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear
- 45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:

	AGTTCCTGGT ACTTTATTAC CAAGGTGACC ATCGGAACCA GGAATGACAT TACTCACTAT	60
50	CAGAATTGAG AAAATTGGTT TGAAAGATGC TGGGCAGTGC ATGATCCCT ATATTACAGT	120
	TAGTGTAAG GATCTGAATG GCATAGACTT AACTCCTGTG GAAATAGCT CTGTGGCTTC	180
55	AAGAAAAGAA GATACATATG TTCAITTTAA TGTGGACATT GATCTTAAAG AGCATGTTGA	240
	AAAATTAACC AAAGGTGCAG CTATCTTCTT TGAATTCAAA GATTAAGG CTAAAAAAG	300
	GTTTACCAGC ACCAAGTGTT TTGCTTTCAT GGAGATGGAT AAAATTAAC CTGGGCCAAT	360
60	TGTAATAGAA CTATACAAGA AACCCACTGA CTTTAAAGA AAAAATTCG AATTATTGAC	420

	CAAGAAACCA CTTTATCTTC ATCTACATCA AACTTTGCAC AAGGAATGAT CCTGACATGA	480
5	TGAACCTGGA ACTTCTGTGA ATTTTACCAC TCAGTAGAAA CCATCATAGC TCTGTGTAGC	540
	ATATTCACCC TTCAACAGGC AGGAAGCAAG CCGTACCCAG ACCAGTAGGC CGGACGGAGT	600
	CAATNGCAAA GCTGTACCAC AGAATTCAGA GTCCAGCACA TCACACTGAC GTATAGGACT	660
10	CCTTGGGATA CAGGTTTATF GTAGATTTTG AAACATGTTT TTACTTTTCT ATTAATTGTG	720
	CAATTAATAG TCTATTTTCT AATTTACCAC TACTCCTACC CTGCTTCCTG GAACAATACT	780
15	GTTGTGGGTA GGATGTGCTC ATCTTCAGAC TTAATACAGC AATAAGAATG TGCTAGAGTT	840
	TACACATCTG TTCACTTTTG CTCCAATATG CTCTTTTGAC TTAACGTCAA GCTTTGGGTT	900
	GATGTGGGTA GGGTAGTGTC AAACGTCTTT GAGAGGAATG GGACCAGTTC TGCTGCCTAA	960
20	GAAGGTCTGT CTGGATGTTT ATAGGCAGCA CCTCTGAAGT GGCCTAAATT CACCCTGATC	1020
	TGATAGTTTT CTGCTTAGA AAGTGTGCCT TGGCCAGATC AGTATCCCAC ATGGGAGTGT	1080
25	TCCCTAGGTT GTAGCTGTGA TTGTTTCCAG ATGACCAGAT TGTTTTCTG AAAATGAGCA	1140
	TATTTTATAGT CATGTGATTT AGCTGTTCTT CTACATCACA TTGTTACTCT TTCTGATGAT	1200
	GATTCTAGGG TTAACATTGG AACCATCTCA AAATAATTAC AAAGTTTATG ATGGGTTTAC	1260
30	AATGTCTTCT AAACAATGTA ATCTAAAAAT AATTGAGTCA GATGCTAACG AGATACTGCA	1320
	GGCATAACTG CTGTTTTTCT GACAACTGAT TGTGAAACCT TAAAACCTGC ATACCTCTTC	1380
35	TTACAGTGAG GAGTATGCAA AATCTGGAAA GATATTCTAT TTTTTTTATA TAGGTAGATA	1440
	GGATCGCCAT TTATTTCCCTA TTTAGATATA CTGACATTCA TCCATATGAA AATATGCAGG	1500
	TCATTAGCTT ACTATAATTT ACTTTTGACT TAATGGGGCA TAAATAAAAC TTTCATAGTA	1560
40	CACATGAGGT GGATATTTGA TACACAGAAC ATTTGCGGTG GGCTTTCTGT GGGTTAGATG	1620
	TAAAGCCAC ATATTTTAAT ATTCACATTT TTAATGAGC AATGCATGAG GGAATGCAG	1680
45	TGTCAGTACC TGGCCTATTT TTAAACTAGT GTAATCACCC TAGTCATACC ATTCAGTATG	1740
	TTTGCTTTTT AAAATAAGTA ACCACAATTA AGTTGTTGTA GCCCTTGCAC TTCAAGAGAT	1800
	CTAGTCTTTA CTTTCAGTTG TCTGTTAGGT CCATTTCTGT TACTAGACGG ATGTTAATAA	1860
50	AAACTATGCG AGCCTGAATG AATTCTCAGC CAAATTTAGT CTTGTCTCTC ATCTTGATTG	1920
	GATTAATTC AAATTTCTAAA ATGATTCAGT CCACAATAGC TCTAGGGGAT GAAGAATTTG	1980
55	CCTTACTTTG CCCAGTTCCT AAGACTGTGA GTTGTCAAAT CCCTAGACTG TAAGCTCTTC	2040
	AAGGAGCAAG AGGCGCATTT TCTCCGTGTC ATGTAATTTT TCTAAGGTGT TTGGCAGCAC	2100
	TCTGTACCCT GTGGAGTACT CAGTACCTTT TGTTTGATGT TGCTGACAAG ACCTGAAAAA	2160
60	AAATCCCTTA AAAAAAAAC CCATTAAAGT GTAGCAAAAC CGAAAAAAA AAAANAAAAA	2220

ACTCGAGACG GGCCCCG

2237

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(2) INFORMATION FOR SEQ ID NO: 105:

10

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1822 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 105:

	GGTCGACCCA CGCGTCCGGA ATTTTCGTAG CAATAAGTTT GTGCATGTAT AGTAATTGTC	60
	ATTAGCAAGG TTGTAACCTC TGCCTCTTGG GTTCAAGTGA TTCTCGTGCC CCAGCCTCCC	120
20	GAGTAGCTGG GACTACAGGC ACGTGCCACC ACGCCCAGCT AATTTTATA TTTTATAGTAG	180
	AGACGGGGTT TTGCTGTGTT GGCCAGGCTG GTCTCAAACCT CCTGACCTCA AGTAATCCAC	240
25	CTGGCCTGCT CTTTTCATGT CTTAACATGG CATGTCMTT AGTTTCATTA TTTTCCTACT	300
	CCTTGATATGT CAAGAAATTA CATTTTGCAT GTCTTATGGA GATGCTGTTA ATTGCTTCAG	360
	TGAGTGCTTT TCTAATCTGC AGACCATTTA CATTTCTGT TTGCAGCATG CTGTGTGCAA	420
30	ACACTCAGTA ATTTGGAGTA TTCAATTATT TGTTAGGGCT CTTCTATTTT CCAAATGTGC	480
	TGAATGTCTT ATTGATGGGA TTTTCAGATC TTTTCATGAG AACTGGAAAT GTAGCTGGGT	540
35	GGCACCTACC TAGGTTGCTA CGTAGTGAGT AGACTTTCTC TTGGGTATAG TAAGCCTCAG	600
	ACAGCTTTCA CTTTATCTA CTTTACTTGT GGAAATAAAA CAGTCATTTT GTTCTGAAAG	660
	AATAAGATAG CTTTCTGTAG AGAAGGAATT CCTACCTCTA AAAGCTGCCT TGAGAACTCA	720
40	GAACTGGCAG TTTTCTGAGG TGATTTTAA ATTTTCAGTAT TAGGGAGAGT CCAGCATTTG	780
	CTGACACAGA TTCTACATAA CTAATGTATG ATAGCAAATG CAAAACCTATT ATAATGTGGT	840
45	GTATCTTGCG CATAACAGG TTAGAACAAG TAGACTCTGG CAGCAGATCT CCAGAGACCC	900
	AAGTTTAGGT TCTCATAGTG TATTTGAAGT AGTTATACTC CTGGCTTAAG TAGTTTAGTG	960
	CCTGGGAGAA TCCATTACTG AAAAGCATTT AACTTAAAAA AAAAAAAAAA AAAAAAAAAA	1020
50	AAACCTCGTG CCGAATTCGG CACGAGCTAA CCCAGAAACA TCCAATTCTC AAACCTGAAGC	1080
	TGCACTCTC GCCTCCAGCA TGAAAGTCTC TGCCGCCCTT CTGTGCCTGC TGCTCATAGC	1140
55	AGCCACCTTC ATTCCCCAAG GGCTCGCTCA GCCAGATGCA ATCAATGCCC CAGTCACCTG	1200
	CTGYTATAAC TTCACCAATA GGAAGATCTC AGTGCAGAGG CTCCCGAGCT ATAGAAGAAT	1260
60	CACCAGCAGC AAGTGTCCTA AAGAAGCTGT GATCTTCAAG ACCATTCTGG CCAAGGAGAT	1320

CTGTGCTGAC CCCAAGCAGA AGTGGGTTCA GGATTCCATG GACCACCTGG ACAAGCAAAC 1380
 CCAAACCTCCG AAGACTTGAA CACTCACTCC ACAACCCAAG AATCTGCAGC TAACTTATTT 1440
 5 TCCCCTAGCT TTCCCCAGAC ACCCTGTTTT ATTTTATTAT AATGAATTTT GTTTGTTGAT 1500
 GTGAAACATT ATGCCTTAAG TAATGTAAAT TCTTATTTAA GTTATTGATG TTTTAAGTTT 1560
 10 ATCTTTCATG GTACTAGTGT TTTTATAGATA CAGAGACTTG GGGAAATTGC TTTTCCTCTT 1620
 GAACCACAGT TCTACCCCTG GGATGTTTTG AGGGTCTTTG CAAGAATCAT TAATACAAAG 1680
 AATTTTTTTT AACATTCCAA TGCATTGCTA AAATATTATT GTGGAAATGA ATATTTTGTA 1740
 15 ACTATTACAC CAAATAAATA TATTTTGTGA CAAAAAAAAA AAAAAAAAAA AAAAAAAAAA 1800
 AAGSGGCCGC TCGAATTAAG CC 1822

20

(2) INFORMATION FOR SEQ ID NO: 106:

(i) SEQUENCE CHARACTERISTICS:
 25 (A) LENGTH: 1712 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 106:

CGTGCCCCAG CCTCCCGAGT AGCTGGRAC ACAGGCACGT SCCACCACGC CCAGCTAATT 60
 TTWATATTTT WAGTAGAGAC GGGGTTTTSC TGKTTGGCC AGGCTGGTCT CAAACTCCTG 120
 35 ACCTCAAGTA ATCCACCTGG CCTGCTCTTT TCATGTCTTA ACATGGCATG TCTTTTAGTT 180
 TCATTATTTT CCTACTCCTT GTATGTCAAG AAATTACATT TTGCATGTCT TATGGAGATG 240
 40 CTGTTAATTG CTTCACTGAG TGCTTTTCTA ATCTGCAGAC CATTTACATT TCCTGTTTGC 300
 AGCATGCTGT GTGCAACAC TCAGTAATTT GGAGTATTCA ATTATTTGTT AGGGCTCTTC 360
 CTATTTCCAA ATGTGCTGAA TTGTCTATTG ATGGGATTTT CAGATCTTTT CATGAGAACT 420
 45 GGAAATGTAG CTGGGTGGCA CCTACCTAGG TTGCTACGTA GTGAGTAGAC TTTCTCTTGG 480
 GTATAGTAAG CCTCAGACAG CTTTCACTTT TATCTACTTT ACTTGTGGAA ATAAAACAGT 540
 50 CATTTTCTTC TGAAAGAATA AGATAGCTTT CTGTAGAGAA GGAAATTCCTA CCTCTAAAAG 600
 CTGCCTTGAG AACTCAGAAC TGGCAGTTT CTGAGGTGAT TTTTAAATTT CAGTATTAGG 660
 GAGAGTCCAG CATTTGCTGA CACAGATTCT ACATAACTAA TGTATGATAG CAAATGCAAA 720
 55 ACTATTATAA TGTGGTGTAT CTTGCGCATA CACAGGTTAG AACAAGTAGA CTCGTCAGC 780
 AGATCTCCAG AGACCCAAGT TTAGGTTCTC ATAGTGATT TGAAGTAGTT ATACTCCTGG 840
 60 CTTAAGTAGT TTAGTGCTG GGAGAATCCA TTAAGTAAAA GCATTTAACT TAAAAAATAA 900

250

AAAAAAAAAA AAAAAAAAAAC CTCGTGCCGA ATTCGGCACG AGCAGAAACA TCCAATTCTC 960
 AACTGAAGC TCGCACTCTC GCCTCCAGCA TGAAAGTCTC TGCCGCCCTT CTGTGCCTGC 1020
 5 TGCTCATAGC AGCCACCTTC ATTCCCCAAG GGCTCGCTCA GCCAGATGCA ATCAATGCCC 1080
 CAGTCACCTG CTGYTATAAC TTCACCAATA GGAAGATCTC AGTGCAGAGG CTCGCGAGCT 1140
 10 ATAGAAGAAT CACCAGCAGC AAGTGTCCCA AAGAAGCTGT GATCTTCAAG ACCATTGTGG 1200
 CCAAGGAGAT CTGTGCTGAC CCAAGCAGA AGTGGGTTC AAGATTCCATG GACCACCTGG 1260
 ACAAGCAAAC CCAAACCTCCG AAGACTTGAA CACTCACTCC ACAACCCAAG AATCTGCAGC 1320
 15 TAACCTATTT TCCCCTAGCT TTCCCAGAC ACCCTGTTTT ATTTTATTAT AATGAATTTT 1380
 GTTTGTTGAT GTGAAACATT ATGCCTTAAG TAATGTTAAT TCTTATTTAA GTTATTGATG 1440
 20 TTTTAAGTTT ATCTTTCATG GTACTAGTGT TTTTGTAGATA CAGAGACTTG GGGAAATTGC 1500
 TTTTCCTCTT GAACCACAGT TCTACCCCTG GGATGTTTTG AGGGTCTTTG CAAGAATCAT 1560
 TAATACAAAG AATTTTTTTT AACATTCCAA TGCATTGCTA AAATATTATT GTGGAAATGA 1620
 25 ATATTTTGTA ACTATTACAC CAAATAAATA TATTTTGTGA CAAAAAAAAA AAAAAAAAAA 1680
 AAAAAAAAAA AAGSGGCCGC TCGAATTAAG CC 1712
 30

(2) INFORMATION FOR SEQ ID NO: 107:

- 35 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1969 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear
 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 107:

CCCCTCCTTC CCCTYGCCAC CTAAGAACC CTCCTCCGAG GTGCCCAGC AGCCGTCTGC 60
 45 CCAGCCACTC CCTGGGAGTC CCCCCAGAAG AGCCTATTAC ATCTACTCCG GGGGCGAGAA 120
 GATCCCCCTG GTGTTGAGCC GGCCCTCTC CTCCAACGTG GCCACTCTTC AGCATCTCTG 180
 TCGGAAGACC GTCAACGGCC ACCTGGACTC CTATGAGAAA GTCACCCAGC TGCCGGGGCC 240
 50 CATTCGGRAG TTCTTGACC AGTACGATGC CCCGTTTTAA GGGGTAAAGG GCGCAAAGG 300
 CATGGGTCGG GAGAGGGGAC GCAGGCCCTC CTCCTCCGTG GCACATGGCA CAAGCACAAG 360
 55 AAGCCAACCA GGAGAGAGTC CTGTAGCTCT GGGGGGAAAG AGGGCGGACA GGCCCCCTCC 420
 TCTGCCCTCT CCCTGCAGAA TGTGGCAGGC GGACCTGGAA TGTGTTGGAG GGAAGGGGGA 480
 GTACCACCTG AGTCTCCAGC TTCTCCGAG ACCCAGCTGT CCTGGTGGGA CGATAGCAAC 540
 60

	CACAAGTGGGA	TTCTCCTTCA	ATTCCTCAGC	TTCCCCCTCTG	CCTCCAAACA	GGGGACACTT	600
	CGGGAATGCT	GAAYTAATGA	GAAGTGCCAG	GGAATCTTCA	AACTTTCCAA	CGGAACCTGT	660
5	TTGCTCTTTG	ATTTGGTTTA	AACCTGAGCT	GGTTGTGGAG	CCTGGGAAAG	GTGGAAGAGA	720
	GAGAGGTCCT	GAGGGCCCCA	GGGSTGCGGG	CTGGCGAAGG	AAATGGTCAC	ACCCCCCGCC	780
10	CACCCCAGGC	GAGGATCCTG	GTGACATGCT	CCTCTCCCTG	GCTCCGGGGA	GAAGGGCTTG	840
	GGGTGACCTG	AAGGGAACCA	TCCTGGTGCC	CCACATCCTC	TCCTCCGGGN	ACAGTCACCG	900
	AAAACACAGG	TTCCAAAGTC	TACCTGGTGC	CTGAGAGCCC	AGGGCCCTTC	CTCCGTTTTA	960
15	AGGGGGAAGC	AACATTTGGA	GGGGACGGAT	GGGCTGGTCA	GCTGGTCTCC	TTTTCTCTACT	1020
	CATACTATAC	CTTCCTGTAC	CTGGGTGGAT	GGAGCGGGAG	GATGGAGGAG	ACGGGACATC	1080
	TTTCACCTCA	GGCTCCTGGT	AGAGAAGACA	GGGGATTCTA	CTCTGTGCCT	CCTGACTATG	1140
20	TCTGGCTAAG	AGATTCGCCT	TAAATGCTCC	CTGTCCCATG	GAGAGGGACC	CAGCATAGGA	1200
	AAGCCACATA	CTCAGCCTGG	ATGGGTGGAG	AGGCTGAGGG	ACTCACTGGA	GGGCACCAAG	1260
25	CCAGCCCACA	GCCAGGGAAG	TGGGGAGGGG	GGGCGGAAAC	CCATGCCTCC	CAGCTGAGCA	1320
	CTGGGAATGT	CAGCCCAGTA	AGTATTGGCC	AGTCAGGCGC	CTCGTGGTCA	GAGCAGAGCC	1380
30	ACCAGGTCCC	ACTGCCCCGA	GCCCTGCACA	GCCCTCCCTC	CTGCCTGGGT	GGGGGAGGCT	1440
	GGAGGTCATT	GGAGAGGCTG	GACTGCTGCC	ACCCCGGGTG	CTCCCGCTCT	GCCATAGCAC	1500
	TGATCAGTGA	CAATTTACAG	GAATGTAGCA	GCGATGGAAT	TACCTGGAAC	ATTTTTTGTT	1560
35	TTTGTTTTTG	TTTTTGTTTT	TGTGGGGGGG	GGCAACTAAA	CAAACACAAA	GTATTCGTGTG	1620
	TCAGGTATTG	GGCTGGACAG	GGCAGTTGTG	TGTTGGGGTG	GTPTTTTTTCT	CTATTTTTTTT	1680
40	GTTTGTTTCT	TGTTTTTTTAA	TAATGTTTAC	AATCTGCCTC	AATCACTCTG	TCTTTTATAA	1740
	AGATTCCACC	TCCAGTCCTC	TCTCCTCCCC	CCTACTCAGG	CCCTTGAGGC	TATTAGGAGA	1800
	TGCTTGAAGA	ACTCAACAAA	ATCCCAATCC	AAGTCAAAC	TTGCACATAT	TTATATTTAT	1860
45	ATTCAGAAAA	GAAACATTTT	AGTAATTTTAT	AATAAAGAGC	ACTATTTTTTT	AATGAAAAAA	1920
	AAAAAAAAAA	AAAAAAAAAA	CGACGCTGGT	GACCGGAATY	CGACGTACG		1960

50

(2) INFORMATION FOR SEQ ID NO: 108:

55 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1734 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 108:

	CGGGTCCCAA GCCTGTGCCT GAGCCTGAGC CTGAGCCTGA GCCCAGCCG GGAGCCGGTC	60
5	GCGGGGGCTC CGGGCTGTGG GACCGCTGGG CCCCCAGCGA TGGCGACCCT GTGGGGAGGC	120
	CTTCTTCGGC TTGGCTCCTT GCTCAGCCTG TCGTGCCTGG CGCTTTCCTT GCTGCTGCTG	180
	GCGCATGTNC AGACGCCGCC AAGAATTTTCG AGGATGTCAG ATGTAAATGT ATCTGCCCTC	240
10	CCTATAAAGA AAATTCTGGG CATATTTATA ATAAGAACAT ATCTCAGAAA GATTGTGATT	300
	GCCTTCATGT TGTGGAGCCC ATGCCTGTGC GGGGGCCTGA TGTAAGCA TACTGTCTAC	360
15	GCTGTGAATG CAAATATGAA GAAAGAAGCT CTGTCACAA CAAGGTTACC ATTATAATTT	420
	ATCTCTCCAT TTTGGGCCCTT CTACTTCTGT ACATGGTATA TCTTACTCTG GTTGAGCCCA	480
	TACTGAAGAG GCGCCTCTTT GGACATGCAC AGTTGATACA GAGTGATGAT GATATTGGGG	540
20	ATCACCAGCC TTTTGCAAAT GCACACGATG TGCTAGCCCG CTCCCGCAGT CGAGCCAACG	600
	TGCTGAACAA GGTAGAATAT GCACAGCAGC GCTGGAAGCT TCAAGTCCAA GAGCAGCGAA	660
25	AGTCTGTCTT TGACCGGCAT GTTGTCTCA GCTAATTGGG GAATTGAATT CAAGGTGACT	720
	AGAAAGAAAC AGGCAGACAA CTGGGAAAGA ACTGACTGGG NTTTTGCTGG GTTTCATTTT	780
	AATACCTTGT TGATTTTACC AACTGTTGCT GGAAGATTCA AACTGGAAG CAAAACTTG	840
30	CTTGATTTTT TTTTCTTGT AACGTAATAA TAGAGACATT TTTAAAAGCA CACAGCTCAA	900
	AGTCAGCCAA TAAGTCTTTT CCTATTTGTG ACTTTTACTA ATAAAAATAA ATCTGCCTGT	960
35	AAATTATCTT GAAGTCCTTT ACCTGGAACA AGCACTCTCT TTTTCACCAC ATAGTTTAA	1020
	CTTGACTTTC AAGATAATTT TCAGGGTTTT TGTGTGTGT GTTTTTGTG TGTGTGTGTT	1080
	GGTGGGAGAG GGGAGGGATG CCTGGGAAGT GGTAAACAAC TTTTTTCAAG TCACTTTACT	1140
40	AAACAACTT TTGTAAATAG ACCTTACCTT CTATTTTCGA GTTTCATTTA TATTTTGCAG	1200
	TGTAGCCAGC CTCATCAAAG AGCTGACTTA CTCATTTGAC TTTTGCACTG ACTGTATTAT	1260
45	CTGGGTATCT GCTGTGTCTG CACTTCATGG TAAACGGGAT CTAAAATGCC TGGTGGCTTT	1320
	TCACAAAAAG CAGATTTTCT TCATGTACTG TGATGTCTGA TGCAATGCAT CCTAGAACAA	1380
	ACTGGCCATT TGCTAGTTTA CTCTAAAGAC TAAACATAGT CTTGGTGTGT GTGGTCTTAC	1440
50	TCATCTTCTA GTACCTTTAA GGACAAATCC TAAGGACTTG GACACTTGCA ATAAAGAAAT	1500
	TTTATTTTAA ACCCAAGCCT CCCTGGATTG ATAATATATA CACATTTGTC AGCATTTCCG	1560
55	GTCGTGGTGA GAGGCAGCTG TTTGAGCTCC AATGTGTGCA GCTTTGAACT AGGCCTGGGG	1620
	TTGTGGGTGC CTCTTCTGAA AGGTCTAACC ATTATTGGAT AACTGGCTTT TTTCTTCTC	1680
60	TTTGGAATGT AACAATAAAA ATAATTTTGG AAACATCAAA AAAAAAAAAA AAAA	1734

(2) INFORMATION FOR SEQ ID NO: 109:

5

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2003 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 109:

	CGCAGGGGGC GCGCGGCCCG GGGACTCGCA TTCCCCGGTT CCCCCCTCCAC CCCACGCGGC	60
15	CTGGACCATG GACGCCAGAT GGTGGGCACT GGTGGTGCTG GCTGCGTTCC CCTCCCTAGG	120
	GGCAGGTGGG GAGACTCCCG AAGCCCCTCC GGAGTCATGG ACCCAGCTAT GGTTCCTCCG	180
20	ATTTGTGGTG AATGCTGCTG GCTATGCCAG NTTTATGGTA CCTGGCTACC TCCTGGTGCA	240
	GTACTTCAGG CGGAAGAAGT ACCTGGAGAC CGGTAGGGGC CTCTGCTTTC CCCTGGTGAA	300
	AGCTTGTGTG TTTGGCAATG AGCCCAAGGC CTCTGATGAG GTTCCCCTGG CGCCCCGAAC	360
25	AGAGCGGGCA GAGACCACCC CGATGTGGCA GGCCCTGAAG CTGCTCTTCT GTGCCACAGG	420
	GCTCCAGGTG TCTTATCTGA CTTGGGGTGT GCTGCAGGAA AGAGTGATGA CCCGCAGCTA	480
	TGGGGCCACA GCCACATCAC CGGGTGAGCG CTTTACGGAC TCGCAGTTCC TGGTGCTAAT	540
30	GAACCGAGTG CTGGCACTGA TTGTGGCTGG CCTCTCCTGT GTTCTCTGCA AGCAGCCCCG	600
	GCATGGGGCA CCCATGTACC GGTACTCCTT TGCCAGCCTG TCCAATGTGC TTAGCAGCTG	660
35	GTGCCAATAC GAAGCTCTTA AGTTCGTCAG CTCCCCACC CAGGTGCTGG CCAAGGCCTC	720
	TAAGGTGATC CCTGTGATGC TGATGGGAAA GCTTGTGTCT CGGCGCANTA ACGAACAATG	780
	GGAGTACCTG ACAGCCACCC TCATCTCCAT TGGGGTCAGC ATGTTTCTGC TATCCAGCGG	840
40	ACCAGAGCCC CGCAGCTCCC CAGCCACCAC ACTCTCAGGC CTCATCTTAC TGGCAGGTTA	900
	TATTGCTTTT GACAGCTTCA CCTCAAATG GCAGGATGCC TGTTTGCCTA TAAGATGTCA	960
45	TCGGTGAGCA TGATGTTTGG GGTCAATTTT TTCTCCTGCC TCTTACAGT GGGSTCACTG	1020
	CTAGNAACAG GGGGGMCTA CTGGAGGGAA CCCGCTTCAT CGGGCGACAC AGTGAGTTTG	1080
	CTGCCCATGC CCTGCTACTC TCCATCTGCT CCGCATGTGG CCAGCTCTTC ATCTTTTACA	1140
50	CCATTGGGCA GTTTGGGGCT GCCGTCTTCA CCATCATCAT GACCCTCCGC CAGGCCTTTG	1200
	CCATCCTTCT TTCCTGCCTT CTCTATGGCC AACTGTGAC TGTGCTGGGA CGGCTGGGGG	1260
55	TGGCTGTGGT CTTTGCTGCC CTCCTGCTCA GAGTCTACCT CAGGGGCGCT CTAAAGCAAC	1320
	GGGGAAGAA GGCTGTGCCT GTTGAGTCTC CTGTGCAGAA GCTTTGAGGG TGGAAAGGGC	1380
60	CTGAGGGGTG AAGTGAAATA GGACCCTCCC ACCATCCCCT TCTGCTGTAA CCTCTGAGGG	1440

AGCTGGCTGA AAGGGCAAAA TGCAGGTGTT TTCTCAGTAT CACAGACCAG CTCTGCAGCA 1500
 GGGGATTGGG GAGCCCAGGA GGCAGCCTTC CCTTTTGCCT TAAGTCACCC ATCTTCCAGT 1560
 5 AAGCAGTTTA TTCTGAGCCC CGGGGGTAGA CAGTCCTCAG TGAGGGGTTT TGGGGAGTTT 1620
 GGGGTCAAGA GAGCATAGGT AGGTTCACCA GTTACTCTTC CCACAAGTTC CCTTAAGTCT 1680
 TGCCCTAGCT GTGCTCTGCC ACCTTCCAGA CTCAC'TCCCC TCTGCAAATA CCTGCATTTT 1740
 10 TTACCCTGGT GAGAAAAGCA CAAGCGGTGT AGGCTCCAAT GCTGCTTTCC CAGGAGGGTG 1800
 AAGATGGTGC TGTGCTGAGG AAAGGGGATG CAGAGCCCTG CCCAGCACCA CCACCTCCTA 1860
 15 TGCTCCTGGA TCCCTAGGCT CTGTTCCATG AGCCTGTTGC AGGTTTGGT ACTTTAGAAA 1920
 TGTAAC'TTTT TGCTCTTATA ATTTTATTTT ATTAAATTA ATTACTGCAA AAAAAAAAAA 1980
 AAAAAAATCG GGGGGGGGCC CGN 2003
 20

25 (2) INFORMATION FOR SEQ ID NO: 110:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1320 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 30 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 110:

GCTGAGCTGC CTTGAGGTGC AGTGTGGGG ATCCAGAGCC ATGTGGGACC TGCTACTACT 60
 35 GGGCCTGATT GGGGGCCTGA CTCTCTTACT GCTGCTGACG CTGCTGGCCT TTGCCGGGTA 120
 CTCAGGGCTA CTGGCTGGGG TGGAAGTGAG TGCTGGGTCA CCCCCATCC GCAACGTCAC 180
 40 TGTGGCCTAC AAGTTCACCA TGGGGCTCTA TGGTGAGACT GGGCGGCTTT TCACTGAGAG 240
 CTGCAGCATC TCTCCCAAGC TCCGCTCCAT CGCTGTCTAC TATGACAACC CCCACATGGT 300
 GCCCCCTGAT AAGTGCCGAT GTGCCGTGGG CAGCATCCTG AGTGAAGGTG AGGAATCGCC 360
 45 CTCCCCTGAG CTCATCGACC TCTACCAGAA ATTTGGCTTC AAGGTGTTCT CCTTCCCGGC 420
 ACCCAGCCAT GTGGTGACAG CCACCTTCCC CTACACCACC ATTCTGTCCA TCTGGCTGGC 480
 50 TACCCGCCGT GTCCATCCTG CCTTGACAC CTACATCAAG GAGCGGAAGC TGTGTGCCTA 540
 TCCTCGGCTG GAGATCTACC AGGAAGACCA GATCCATTTT ATGTGCCCAC TGGCAGGCA 600
 GGGAGACTTC TATGTGCCTG AGATGAAGGA GACAGACTGG AATGGGGG GCTTGTGGA 660
 55 GGCCATTGAC ACCCAGGTGG ATGGCACAGG AGCTGACACA ATGAGTGACA CGAGTTCTGT 720
 AAGCTTGGA GTGAGCCCTG GCAGCCGGGA GACTTCAGCT GCCACACTGT CACCTGGGGC 780
 60 GAGCAGCCGT GGCTGGGATG ACGGTGACAC CCGCAGCGAG CACAGCTACA GCGAGTCAGG 840

255

5 TGCCAGCGGC TCCTCTTTTG AGGAGCTGGA YTTGGAGGGC GAGGGGCCCT TAGGGGAGTC 900
 ACGGCTGGAC CCTGGGACTK AGCCCCTGGG GACTACCAAG TGGCTCTGGG AGCCCACTGC 960
 CCCTGAGAAG GGCAAGGAGT AACCCATGGC CTGCACCCTC CCTGCAGTGC AGTTGCTGAG 1020
 GAACTGAGCA GACTCTCCAG CAGACTCTCC AGCCCTCTTC CTCCTTCCTC TGGGGGAGGA 1080
 10 GGGGTTCCTG AGGGACCTGA CTTCCCCTGC TCCAGGCCTC TTGCTAAGCC TTCTCCTCAC 1140
 TGCCCTTTAG GCTCCCAGGG CCAGAGGAGC CAGGGACTAT TTTCTGCAAC CAGCCCCCAG 1200
 GGCTGCCNCC CCGTTGTGT CTTTPTTTCA GACTCACAGT GGAGCTTCCA GGACCCAGAA 1260
 15 TAAAGCCAAT GATTACTTG TTTCAAAAAA AAAAWAAAAA AAAAAAAAAA AAAAAAAAAA 1320

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(2) INFORMATION FOR SEQ ID NO: 111:

25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1962 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 111:

30 CGGACCCCTT CCTCCTCCTC NAAGCATGTC CCACCATTGT GGCAGGGGCT GGGGANACAG 60
 TCACCTGATG CGGGGACCAC GGCCACTCCA CCTCGSTGGC GCTGTCACTG GGCAGCACTG 120
 35 GCTGGGCCTG CACTGAGGTC CCTGCTGGGG CAGTTCTTCC AGAATTATCT TCAGAGGGGG 180
 CCTCCAGCTC CCTGGTACCC TCAGGGGCCC GTGTGGCTGG AAGCAGGGAA GGGGCACCCT 240
 40 CGGAGCTTCC TGTCTCCTCG CTCTCTCCTC GAGGGACCCC AGATAGCTCA GGACCACCAG 300
 TTGCCTCCCC CACCTCTCTT GCCTCAACCA GAGTGAAGG TGATGGGGAT GCTAGGTTCC 360
 TCTCCCTGGG AGTGGGCAGA GTCTCAGTAG GTGGTCCATG GACCCTTGGA GGCCTGGAAG 420
 45 CTTCTGACTC TCCATCAGGA AGTGGTGATG CACCAGGCTG CAGGACTGCC CTTGCTGGCG 480
 CCTGGGAGAG TGA CTCTCC TGGGCTGCTG GCTCAGTGGG GAGAGAGGCC TCAGGGCCCG 540
 50 GGCTGCTGAG CTCGCTGGGC CATGCCACA GAGCCTCATC CTCACCTCC TCCTCTCTTT 600
 CTTCTCTCTC TTTCTCTTCT TCATCTTCAT ATTTCTCTTC TTCTTCCAAT GCCTTACCTT 660
 CCTCTTYTGR AAACCCCGTG GCGGTACCA TGGATTGTGT TTCAAATCTT AGGAGCGTCC 720
 55 TAGGGGCCTC TGCTGGGTCT TCTGGAGTGG AGCTTCCAGC TTCTCTCTCT TCCATGATGG 780
 GGATGGAGTA RATGGCCCCA CGGGATTAC TCTCTGTGGC TTCTCTAGGG AGCTGCAGTT 840
 60 CCTCCAGGCT CTCTGTCACT GTGACRATAG CCTCTAGTCC ATCAAAAGCT GGCTTGAGGG 900

256

CTGGGTTGGA GGCCTCAGGG ATGGCAGAAG GCTGGGCCGA GTCTCGGAAG CAGTARACGT 960
 TGAAGCGGCT GTGCTTATTT GGAAGCCAG TCTGGTTGGG GAAGANGAAG AGAGTCTTGA 1020
 5 CACCAGGCAA GCCCCACCA CAGCGCTGGC TGGGTGTGAC GATGGGGTAG CGCACANTGC 1080
 CATCAGCTAG CCACCTGGGC TGCAGTGGTC CAGGCCACCA TCCCAGGCTG CATAAGTTG 1140
 10 GCCCGTGGTG GCAATCTCTG CACCCCGCTC CTGGCAGTAC GCCCGTGCTT CCTCCAATGT 1200
 CAGCTTCTCT GGAGGGTCAC CCAGGAACAG TTCTCCATTT AGGTCTTCAG CATAACAGTA 1260
 CACATCATAG AGGTCATCCG GGTCCACCAC ACCATAGTTC CGGACCCCGG GGAAGCCATC 1320
 15 CATGTCTCCG TAACAGGCCT CTCGTGGGGT CTGGATGGGA TACCTTTGAC CTTGAMCTCC 1380
 ACAGCGTCGC TGCTGTCATC GATGCCGTGC TGGACCTCAC AGCGATAGAT ACCTGAGTCG 1440
 TTGGGGCGCA GCTCGCTCAG CGCCAGGGGA GACGTCGGTG AGCGACGCTG GGTACGCAGG 1500
 20 CAGTGCCACG CGGAACCGGT AGGCCTCGTT CACCTTGACG CGCACTCCCC GCGCCACCAG 1560
 CACYTCTGCC TCCCGGCCCC GGGACAGGAA AGTCCACTTG ACCCGCGGAG AGCCCAGCAC 1620
 25 AGCCCGGCGG CTCGGCGGTG SCCGCAGGTA GTGGACGTGG CAAGGGATGK TGAGGGCSCC 1680
 GCCGAGCAAC GCCYTGCAGT GCGCGTCTGC CCGCGATGCG CACGCGAAAA GCGCGKTCCT 1740
 CTGAGCTGTC TCCTTCCAGA ACATCTGCTA AAGCTGCAGG AGCCTGGGCC AGGACCAGGG 1800
 30 CTGCCAGCAG GGCAGGAAC AGCTGGGCCA TGCTGCAGGC TACCCAGGGC TGGGGTTGGG 1860
 TCGCGGCACT GCGAAGTTTG TCGCCTCCTC CGGGGGTCTC CTCCGGGTCG ACGGCTCAGT 1920
 35 NCCTGCAGCT GCAGCTGAGA CTGCGGCGGA GACTGCGCGA GC 1962

40 (2) INFORMATION FOR SEQ ID NO: 112:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1785 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

45

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 112:

50 AAGTTTCAGC CAAACTTCGG GCGGCTGAGG CGGCGGCCGA GAGCGGGGCG ACTCSGGGCG 60
 CGGGGAGTCG AGGCATTTGC GCCTGGGCTT CGGAGCGTAC GCGAGGCGCT GAGCCTTTGA 120
 AGCAGGAGGA GGGGAGGAGA GAGTGGGGCT CCTCTATCGG CAGGAGTTTC CCAATGTGGAT 180
 55 CTGCCCAGGC GCGGCGGGCG GCCGAGGAGG CGACCGAGAA GATGCTGCTG CTGCGCCCCG 240
 CTCTGCTGTG GCGCTGCTG GCGCTCTGGC TGTGCTGCGC GATGCTGCTG ATGCATTGCA 300

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GTGTCGAGAT GGCTATGAAC CCTGTGTAAA TGAAGGAATG TGTGTTACCT ACCACAATGG 360
 CACAGGATAC TGCAAATGTC CAGAAGGCTT CTTGGGGGAA TATTGTCAAC ATCGAGACCC 420
 5 CTGTGAGAAG AACCGCTGCC AGAATGGTGG GACTTGTGTG GCCCAGGCCA TGCTGGGGAA 480
 AGCCACGTGC CGATGTGCCT CAGGGTTTAC AGGAGAGGAC TGCCAGTACT CGACATCTCA 540
 10 TCCATGCTTT GTGTCTCGAC CTTGCCTGAA TGGCGGCACA TGCCATATGC TCAGCCGGGA 600
 TACCTATGAG TGCACCTGTC AAGTCGGGTT TACAGGTAAG GAGTGCCAAT GGACCGATGC 660
 CTGCCTGTCT CATCCCTGTG CAAATGGAAG TACCTGTACC ACTGTGGCCA ACCAGTTCTC 720
 15 CTGCAAATGC CTCACAGGCT TCACAGGGCA GAAGTGTGAG ACTGATGTCA ATGAGTGTGA 780
 CATTCCAGGA CACTGCCAGC ATGGTGGCAC CTGCCTCAAC CTGCCTGGTT CCTACCAGTG 840
 20 CCAGTGCCTT CAGGGCTTCA CAGGCCAGTA CTGTGACAGC CTGTATGTGC CCTGTGCACC 900
 CTCGCCTTGT GTCAATGGAG GCANCTGTG GCAGACTGGT GACTTCACTT TTGAGTGCAA 960
 CTGCCTTCCA GAAACAGTGA GAAGAGGAAC AGAGCTCTGG GAAAGAGACA GGAAGTCTG 1020
 25 GAATGGAAAA GAACACGATG AGAATTAGAC ACTGGAAAAT ATGTATGTGT GGTTAATAAA 1080
 GTGCTTTAAA CTGAATTGAC ATTAACAGTR GGTGATCAAC TTTMCTATGT GCTTGTGCTT 1140
 30 TTGCTTTTGA TGGAGTAATT CATTGTTTTT TATCCACCT AAATGCACCC AGCTGCCCTT 1200
 GATTTTCTCT GGGCTACTGG CCTTCACAAC CCTCTCCCAT GTACCTCTC TGACTTTGGG 1260
 GTAACCCTCC CCTAACTTAA AGCTAGAGAA TTCTGAACT GAGGAGGGGA TCCTCTGTTA 1320
 35 ATCAGTGAGC ACTTTTTGAT GAGCTGATAG ATGATATATG AGAGACTATG CGTGGCACAA 1380
 TACTTTGTTA CACTCTTCAC TGATACAAGT GTTCTAGAGT GYACACACAA CCCAAAGATA 1440
 40 GAAATAAAAA GAGGAGCAGT GTCGGGGAGC TTGGGGCCTG GTGTTCCATG GAGAGGGAGA 1500
 AAGGAACAAG CTTGRCCAAT TCATTCAACT CCTTATAAAA ATGATGAGGA GGCTGAAAAC 1560
 CAAGAATTTT GATTGGGAAC AGAATACAAG CAGCTGAAC AGATGAWTTA CTAAGCAACA 1620
 45 AAGATCCTGT TTTTATACAA ATATCCTTAG TACAAAAACA AAARAAGGAA AACTGTAGGG 1680
 GGGAGTAATG TGCTAAGTAA GCAGAATTGC CTCCAAAAGA AGTTGTTTCT AGTTACTCTT 1740
 50 TTCCGGGTNG GGATCTTTAG NTTCCGGTAT TGTGGGTATG GTTCC 1785

(2) INFORMATION FOR SEQ ID NO: 113:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1842 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 113:

5	GGAGCCTCTC TTGCAACTTC TGCCACCGCG GGCCACCGCG GCCGCCTGAT CCCGCAGAGG	60
	AAGTCGCGGC CGTGGAGCGA TGACCCGCGG CGGTCCGGGC GGGCGCCCGG GGCTGCCACA	120
	GCCGCCGCGG CTTCTGCTGC TGCTGCTGCT GCMGCTGTG TTAGTCACCG CGGAGCCGCC	180
10	GAAACCTGCA GGAGTCTACT ATGCAACTGC ATACTGGATG CCTGCTGAAA AGACAGTACA	240
	AGTCAAAAAT GTAATGGACA AGAATGGGGA CGCCTATGGC TTTTACAATA ACTCTGTGAA	300
	AACCACAGGC TGGGGCATCC TGGAGATCAG AGCTGGCTAT GGCTCTCAAA CCCTGAGCAA	360
15	TGAGATCATC ATGTTTGTGG CTGGCTTTTT GGAGGGTAC CTCACTGCCC CACACATGAA	420
	TGACCACTAC ACAAACCTCT ACCCAGAGCT GATCACGAAA CCTTCCATCA TGGATAAAGT	480
20	GCAGGATTTT ATGGAGAAGC AAGATAAGTG GACCCGAAA AATATCAAAG AATACAAGAC	540
	TGATTCATTT TGGAGACATA CAGGCTATGT GATGGCACAA ATAGATGGCC TCTATGTAGG	600
	AGCAAAGAAG AGGGCTATAT TAGAAGGGAC AAAGCCAATG ACCCTGTTCC AGATTCAGTT	660
25	CCTGAATAGT GTTGGAGATC TATTGGATCT GATTCCCTCA CTCTCTCCCA CAAAAACGG	720
	CAGCCTAAAG GTTTTAAAGA GATGGGACAT GGGACATTGC TCCGCTCTTA TCAAGGTTCT	780
30	TCCTGGATTT GAGAACATCC TTTTGTCTCA CTCAAGCTGG TACACGTATG CAGCCATGCT	840
	CAGGATATAT AAACACTGGG ACTTCAACRT CATAGATAAA GATACCAGCA GTAGTCGCCT	900
	CTCTTTCAGC AGTTACCCAG GGTTTTTGA GTCTCTGGAT GATTTTACA TTCTTAGCAG	960
35	TGGATTGATA TTGCTGCAGA CCACAAACAG TGTGTTTAAAT AAAACCCTGC TAAAGCAGTA	1020
	ATACCCGAGA CTCTCCTGTC CTGGCAAAGA GTCCGTGTGG CCAATATGAT GGCAGATAGT	1080
40	GGCAAGAGGT GGCAGACAT CTTTTCAAAA TACAACTCTG GCACCTATAA CAATCAATAC	1140
	ATGGTTCTGG ACCTGAAGAA AGTAAAGCTG AACCACAGTC TTGACAAAGG CACTCTGTAC	1200
	ATTGTGGAGC AAATTCCTAC ATATGTAGAA TATTCTGAAC AAAGTATGT TCTACGAAA	1260
45	GGATATTGGC CCTCCTACAA TGTTCTTTTC CATGAAAAAA TCTACAACTG GAGTGGCTAT	1320
	CCACTGTTAG TTCAGAAGCT GGGCTTGGAC TACTCTTATG ATTTAGCTCC ACGAGCCAAA	1380
50	ATTTTCCGGC GTGACCAAGG GAAAGTGAAT GATACGGCAT CCATGAAATA TATCATGCGA	1440
	TACAACAATT ATAAGAAGGA TCCTTACAGT AGAGGTGACC CCTGTAATAC CATCTGCTGC	1500
	CGTGAGGACC TGAACTCACC TAACCCAAGT CCTGGAGGTT GTTATGACAC AAAGGTGGCA	1560
55	GATATCTACC TAGCATCTCA GTACACATCC TATGCCATAA GTGGTCCCAC AGTACAAGGT	1620
	GGCCTCCCTG TTTTTCGCTG GGACCGTTTC AACAAAATC TACATCAGGG CATGSCAGAG	1680
60	GTCTACAAC TTTGATTTTAT TACCATGAAA CCAATTTTGA AACTTGATAT AAAATGAAGG	1740

AGGGAGATGA CGGACTAGAA GACTGTAAAT AAGATACCAA AGGCACTATT TTAGCTATGT 1800
 TTTTCCCATC AGAATTATGC AATAAAATAT ATTAATTTGT CA 1842

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(2) INFORMATION FOR SEQ ID NO: 114:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1960 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

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(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 114:

GAATTCGGCA CGAGCTTCTC CGCGCCCCAG CCGCCGGCTG CCAGCTTTTC GGGGCCCCGA 60
 20 GTCGCACCCA GCGAAGAGAG CGGGCCCCGG ACAAGCTCGA ACTCCGGCCG CCTCGCCCTT 120
 CCCC GGCTCC GCTCCCTCTG CCCCCTCGGG GTCGCGCGCC CACGATGCTG CAGGGCCCTG 180
 25 GCTCGCTGCT GCTGCTCTTC CTCGCCCTCGC ACTGCTGCCT GGGCTCGGCG CGCGGGCTCT 240
 TCCTCTTTGG CCAGCCCGAC TTCTCCTACA AGCGCAGMAA TTGCAAGCCC ATCCCGGTCA 300
 ACCTGCAGCT GTGCCACGGC ATCGAATACC AGAACATGCG GCTGCCCAAC CTGCTGGGCC 360
 30 ACGAGACCAT GAAGGAGGTG CTGGAGCAGG CCGGCGCTTG GATCCCGCTG GTCATGAAGC 420
 AGTGCCACCC GGACACCAAG AAGTTCTTGT GCTCGCTCTT CCCCCCGTC TGCCTCGATG 480
 35 ACCTAGACGA GACCATCCAG CCATGCCACT CGCTCTGCGT GCAGGTGAAG GACCGCTGCG 540
 CCCC GGTCAT GTCCGCCTTC GGNTTCCCTT GGGCCGACAT GCTTGAGTGC GACCGTTTCC 600
 CCCAGGACAA CGACCTTTGC ATCCCCCTCG CTAGCAGCGA CCACCTCCTG CCAGCCACCG 660
 40 AGGAAGCTCC AAAGGTATGT GAAGCCTGCA AAAATAAAAA TGATGATGAC AACGACATAA 720
 TGGAAACGCT TTGTAAAAAT GATTTTGCAC TGAAAATAAA AGTGAAGGAG ATAACCTACA 780
 45 TCAACCGAGA TACCAAAATC ATCCTGGAGA CCAAGAGCAA GACCATTAC AAGCTGAACG 840
 GTGTGTCCGA AAGGGACCTG AAGAAATCGG TGCTGTGGCT CAAAGACAGC TTGCAGTGCA 900
 CCTGTGAGGA GATGAACGAC ATCAACGCGC CCTATCTGGT CATGGGACAG AAACAGGGTG 960
 50 GGGAGCTGGT GATCACCTCG GTGAAGCGGT GGCAGAAGGG GCAGAGAGAG TTCAAGCGCA 1020
 TCTCCCGCAG CATCCGCAAG CTGCAGTGCT AGTCCCGGCA TCCTGATGGC TCCGACAGCC 1080
 55 CTGCTCCAGA GCACGGCTGA CCATTTCTGC TCCGGGATCT CAGCTCCCCG TCCCCAAGCA 1140
 CACTCCTAGC TGCTCCAGTC TCAGCCTGGG CAGCTTCCCC CTGCCCTTTTG CACGTTTGCA 1200
 TCCCCAGCAT TTCCTGAGTT ATAAGGCCAC AGGAGTGGAT AGCTGTTTTT ACCTAAAGGA 1260
 60

260

AAAGCCCACC CGAATCTTGT AGAAATATTC AACTAATAA AATCATGAAT ATTTTATGA 1320
 AGTTTAAAAA TAAAGCTAGT TTTGAATAGG TGCAACTGTG ACTTGGGTCT 1380
 5 GGTGGTGGT TTTGAGTCAG CTGATTTTCA CTTCCCACTG AGGTTGTCAT 1440
 AACATGCAAA TTGTTTCAAT TTTCTCTGTG GCCCAAACCT GTGGGTCACA AACCTGTGTG 1500
 AGATAAGCT GGCTGTTATC TCAACATCTT CATCAGCTCC AGACTGAGAC TCAGTGTCTA 1560
 10 AGTCTTACAA CAATTCATCA TTTTATACCT TCAATGGGAA CTTAAACTGT TACATGTATC 1620
 ACATTCCAGC TACAATACTT CCATTTATTA GAAGCACATT AACCATTCTT ATAGCATGAT 1680
 15 TTCTTCAAGT AAAAGGCAAA AGATATAAAT TTTATAATTG ACTTGAGTAC TTTAAGCCTT 1740
 GTTTAAAACA TTTCTTACTT AACTTTTGCA AATTAAACCC ATTGTAGCTT ACCTGTAATA 1800
 TACATAGTAG TTTACCTTTA AAAGTTGTAA AAATATTGCT TTAACCAACA CTGTAAATAT 1860
 20 TTCAGATAAA CATTATATTC TTGTATATAA ACTTTACATC CTGTTTTACC TAAAAAATAA 1920
 AAAAAAATAA AAAAAACTCG AGGGGGGCCC GGTACCCAAT 1960
 25

(2) INFORMATION FOR SEQ ID NO: 115:

30 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 536 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 35 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 115:

GTGCTCAGCC CCCGGGGCAC AGYAGGACGT TTGGGGGCCT TCTTTCAGCA GGGGACAGCC 60
 40 CGATTGGGGA CAATGGCGTC TCTTGCCAC ATCTTGGTTT TCTGTGTGGG TCTCCTCACC 120
 ATGGCCAAGG CAGAAAGTCC AAAGGAACAC GACCCGTTC AATTACGACTA CCAGTCCCTG 180
 CAGATCGAGG GCCTCGTCAT CGCCGGGATC CTCTTCATCC TGGGCATCCT CATCGTGCTG 240
 45 AGCAGAAGAT GCCGGTGCAA GTTCAACCAG CAGCAGAGGA CTGGGGAACC CGATGAAGAG 300
 GAGGGAACCT TCCGCAGCTC CATCCGCCGT CTGTCCAMCC GCANGCGGTA GAAACACCTG 360
 50 GAGCGATGGA ATCCGGCCAG GACTCCCCTG GCACCTGACA TCTCCCACGC TCCACCTGCC 420
 CGCCACCGC CCCCTCCGCC GCCCTTCCC CAGCCCTGCC CCCGCAGACT CCCCTGCCG 480
 CCAAGACTTC CAATAAACG TCGTTTCCTC TCGAMAAAAA AAAAAATAAA AAAACT 536
 55

60 (2) INFORMATION FOR SEQ ID NO: 116:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 790 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 116:

GTGGGGAGGG GCGGAGCAA AGCCGCGCCT CTGGGTGGGC GGGTCGGGCC GTCCAGGTCC 60
 CTGACTTGAA CCTTCCCGGT CCCAGCCCT CAACAGGAGG CGCAGAAAAT CTTCAAAGCC 120
 AACCACCCCA TGGACGCAGA AGTTACTAAG GCCAAGCTTC TGGGGTTTGG CTCTGCTCTC 180
 CTGGACAATG TGGACCCCAA CCTGAGAAC TTCGTGGGGG CGGGGATCAT CCAGACTAAA 240
 GCCCTGCAGG TGGGCTGTCT GCTTCGGCTG GAGCCCAATG CCCAGGCCCA GATGTACCGG 300
 CTGACCCTGC GCACCAGCAA GGAGCCCGTC TCCCGTCACC TGTGTGAGCT GCTGGCACAN 360
 AGTTCTGAGC CCTGGACTCT GCGCCGGGGG ATGTGGCCGG CACTGGGCAG CCCCTTGGAC 420
 TGAGGCAGTT TTGGTGGATG GGGGACCTCC ACTGGTGACA GAGAAGACAC CAGGGTTTGG 480
 GGGATGCCTG GGACTTTCCT CCGGCCTTTT GTATTTTAT TTTTGTTCAT CTGCTGCTGT 540
 TTACATTCTG GGGGGTTAGG GGGAGTCCCC CTCCCTCCCT TTCCCCCCA AGCACAGAGG 600
 GGAGAGGGGC CAGGGAAGTG GATGTCTCCT CCCCTCCCAC CCCACCCTGT TGTAGCCCCT 660
 CCTACCCCCT CCCCATCCAG GGGCTGTGTA TTATTGTGAG CGAATAAACA GAGAGACGTT 720
 AACAGCCCCA TGTCTGTGTC CATCACCCAN TGNTAGGTAG TCAAAGAAGT GGGGTGAGGG 780
 CATGCAGAGT 790

(2) INFORMATION FOR SEQ ID NO: 117:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 776 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 117:

CAGCGCTGGA AGCAGCTGAG CCGTGAGGG GTGGGGAGGG GCGGAGCAA AGCCGCGCCT 60
 CTGGGTGGGC GGGTCGGGCC GTCCAGGTCC CTGACTTGAA CCTTCCCGGT CCCAGCCCT 120
 CAACAGGAGG CGCAGAAAAT CTTCAAAGCC AACCACCCCA TGGACGCAGA AGTTACTAAG 180
 GCCAAGCTTC TGGGGTTTGG CTCTGCTCTC CTGGACAATG TGGACCCCAA CCTGAGAAC 240
 TTCGTGGGGG CGGGGATCAT CCAGACTAAA GCCCTGCAGG TGGGCTGTCT GCTTCGGCTG 300
 GAGCCCAATG CCCAGGCCCA GATGTACCGG CTGACCCTGC GCACCAGCAA GGAGCCCGTC 360

TCCCGTCACC TGTGTGAGCT GCTGGCACAG AGTTCTGAGC CCTGGACTCT GCCCCGGGGG 420
 ATGTGGCCCG CACTGGGCAG CCCCTTGGAC TGAGGCAGTT TTGGTGGATG GGGGACCTCC 480
 5 ACTGGTGACA GAGAAGACAC CAGGGTTTGG GGGATGCCTG GGACTTTCCT CCGGCCTTTT 540
 GTATTTTAT TTTTGTTCAT CTGCTGCTGT TTACATTCTG GGGGTTAGG GGGAGTCCCC 600
 10 CTCCCTCCCT TTCCCCCCCCA AGCACAGAGG GGAGAGGGGC CAGGGAAGTG GATGTCTCCT 660
 CCCCTCCCAC CCCACCCTGT TGTAGCCCCT CCTACCCCCT CCCCATCCAG GGGCTGTGTA 720
 TTATTGTGAG CGAATAAACA GAGAGACGCN TAAAAAATAA AAAAAAAT TGAGGG 776
 15

(2) INFORMATION FOR SEQ ID NO: 118:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 453 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

25

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 118:

GGTTCTGACA CCAGATGTTT TCTGCTCCTG GTTAATGTCA GTGAGGGCTG GAAGTTGAAT 60
 30 AAATGAGAAC AGGAGTGGTC TGGGCCCCATG TAAATGATCC TCCCTTGAAA GGAGGAACAG 120
 CTTTCATCAT TTGTCCAGC TAAGCCTTGC ATGCATTATA GATCTGGTGC TAAGCAGTGG 180
 35 GAAAGATCTC ATAAGTAATG TTTTATGTTT TTTCKGTCTC TCYTCTTCKG TTGTCTTGG 240
 CTGTGGGTT GTGTTTGKGG TTGTTAACTG GAAAATTGCT ATAAGCCAGT TGTCYKAAK 300
 TTTWAAAAAC GAATTAGAAA AACCATAAAA TCYTCTGGCC YATGCACATK GTCCCYGTTT 360
 40 TGTGAAAACA TTAAAGGGTA AATAAAAAGG AAGGAGAACA GTCAATAATG TGCATCAAAT 420
 ATATTCTGAG TTCTAGAGAA ATTAATGACC AAG 453
 45

(2) INFORMATION FOR SEQ ID NO: 119:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2016 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 119:

AGGCTGTTCA CAGGCACCCC GAGACAGCGT CCCCCCTCTG GGGGACTCG ATTTGACGTT 60
 60 GCAGGACGCG CGGCTGGAAC CCCAGGCCCG CGCTGCTCAC AGACGGGAC TCCGCCTCCG 120

	TCCTAGAGCG AGAATTTGGGG GAAAGCTGTT ATTTTATAT TAAAATACAT TCAGATGTAA	1980
5	AAAAAAAAA AAAAAAANC'T CGAGGGGGGG CCCCCG	2016
10	(2) INFORMATION FOR SEQ ID NO: 120:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 2136 base pairs	
	(B) TYPE: nucleic acid	
15	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 120:	
20	GGGGACGGAG CCGCTGTCAA CTCTCCAAC'T CAGCTCAGCT GATCGGTTGC CGCCGCCGCC	60
	GCCGCCAGAT TCTGGAGGCG AAGAACGCAA AGCTGAGAAC ATGGACGTTA ATATCGCCCC	120
	ACTCCGCGCC TGGGACGATT TCTTCCGGG TTCCGATCGC TTTGCCCGGC CGGACTTCAG	180
25	GGACATTTCC AAATGGAACA ACCGCGTAGT GAGCAACCTG CTCTATTACC AGACCAACTA	240
	CCTGGTGGTG GCTGCCATGA TGATTTCAT TGTTGGGTTT CTGAGTCCCT TCAACATGAT	300
30	CCTGGGAGGA ATCGTGGTGG TGCTGGTGT CACAGGGTTT GTGTGGGCAG CCCACAATAA	360
	AGACGTCCTT CGCCGGATGA AGAAGCGCTA CCCCACGACG TTCGTTATGG TGGTCATGTT	420
	GGCGAGCTAT TTCCTTATCT CCATGTTTGG AGGAGTCATG GTCTTTGTGT TTGGCATTAC	480
35	TTTTCCTTTG CTGTTGATGT TTATCCATGC ATCGTTGAGA CTTCGGAACC TCAAGAACAA	540
	ACTGGAGAAT AAAATGGAAG GAATAGGTTT GAAGAGGACA CCGATGGGCA TTGTCCTGGA	600
40	TGCCCTAGAA CAGCAGGAAG AAGGCATCAA CAGACTCACT GACTATATCA GCAAAGTGAA	660
	GGAATAAACA TAACTTACCT GAGCTAGGGT TGCAGCAGAA ATTGAGTTGC AGCTTGCCCT	720
	TGTCCAGACC TATKTTCTGC TTGCGTTTTT GAAACAGGAG GTGCACGTAC CACCCAATTA	780
45	TCTATGGCAG CATGCATGTA TAGGCCGAAC TATTATCAGC TCTGATGTTT CAGAGAGAAG	840
	ACCTCAGAAA CCGAAAGAAA ACCACCACC TCCTATTGTG TCTGAAGTTT CACGTGTGTT	900
50	TATGAAATCT AATGGGAAAT GGATCACACG ATTCTTTTAA GGGAAATTAA AAAAAATAAA	960
	GAATTACGGC TTTTACAGCA ACAATACGAT TATCTTATAG GAAAAAATAA ATCATTTGTAA	1020
	AGTATCAAGA CAATACGAGT AAATGAAAAG GCTGTTAAGG TAGATGACAT CATGTGTTAG	1080
55	CCTGTTCCCTA ATCCCCTAGA ATTGTAATGT GTGGGATATA AATTAGTTT TATTATTCTC	1140
	TTAAAAATCA AAGATGATCT CTATCACTTT GCCACCTGTT TGATGTGCAG TGGAAACTGG	1200
60	TTAAGCCAGT TGTTCACTACT TCSTTTACAA ATATAAAGAT AGCTGTTTAG GATATTTTGT	1260

265

TACATTTTGTG TAAATTTTGTG AAATGCTAGT AATGTGTTTTT CACCAGCAAG TATTTGTTGC 1320
 AAACCTTAATG TCATTTTCCT TAAGATGGTT ACAGCTATGT AACCTGTATT ATTCTGGACG 1380
 5 GACTTATTAA AATACAAACA GACAAAAAAT AAAACAAAAC TTGAGTTCTA TTTACCTTGC 1440
 ACATTTTGTG TTGTTACAGT GAAAAAAATG GTCCAAGAAA ATGTTTGCCA TTTTTCATT 1500
 GTTTCGTTTT TAACTGGAAC ATTTAGAAAAG AAGGAAATGA ATGTGCATTT TATTAATTCC 1560
 10 TTAGGGGCAC AAGGAGGACA ATAATAGCTG ATCTTTTGAA ATTTGAAAAA CGTCTTTAGA 1620
 TGACCAAGCA AAAAGACTTT AAAAAATGGT AATGAAAATG GAATGCAGCT ACTGCAGCTA 1680
 15 ATAAAAAATT TTAGATAGCA ATTGTTACAA CCATATGCCT TTATAGCTAG ACATTAGAAT 1740
 TATGATAGCA TGAGTTTATA CATTCCTATTA TTTTCTCTCC CTTTCTCATG TTTTATATAA 1800
 TAGGTAATAA AAAATGTTTT GCCTGCCAAT TGAATGATTT CGTAGCTGAA GTAGAAACAT 1860
 20 TTAGGTTTCT GTAGCATTAA ATTGTGAAGA CAACTGGAGT GGTACTTACT GAAGAACTC 1920
 TCTGTATGTC CTAGAATAAG AAGCAATGAT GTGCTGCTTC TGATTTTCT TGCATTTTAA 1980
 25 ATTCTCAGCC AACCTACAGC CATGATCTTT AGCACAGTGA TATCACCATG ACTTCACAGA 2040
 CATGGTCTAG AATCTGTACC CTTACCCACA TATGAAGAAT AAAATTGATT AAAGGTTAAA 2100
 AAAAAAWAA AAAAAMWAGG GGGGCCCGGT WCCCAG 2136
 30

35 (2) INFORMATION FOR SEQ ID NO: 121:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 219 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 40 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 121:

45 GCCCTAGTAT CTGGGCAGCT GTGCATGGAG ATAGCCAGAG GAAACATTTT TTTTCTTAAT 60
 GRATTGGTGA CCACATTTTG TTGTCTTTCG CTCCTATTAT CCGTGCCTA TTTGCATSCT 120
 GGTTCCTTCT ACAGTAGTTT ATGTAAATGT TGTTCCTTCT TTGTCGTTCT CAGTAGAATT 180
 50 GGTTCGTAA ACGAAACCTG GTCCTGTAAT TTCAGTATA 219

55 (2) INFORMATION FOR SEQ ID NO: 122:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1686 base pairs
 (B) TYPE: nucleic acid
 60 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 122:

5	GCTGGAGATT CACATTTTAC CTGATTGCCT TCATMGCCGG CATGGCCGTC ATTGTGGATA	60
	AACCCTGGTT CTATGACATG AAGAAAGTTT GGGAGGGATA TCCCATACAG AGCACTATCC	120
	CTTCCCAGTA TTGGTACTAC ATGATTGAAC TTTCTTCTA CTGGTCCCTG CTCTTCAGCA	180
10	TTGCCTCTGA TGTCAAGCGA AAGGATTTCA AGGAACAGAT CATCCACCAT GTGRCCACCA	240
	TCATTTCTCAT CAGCTTTTCC TGGTTTGCCA ATTACATCCG AGCTGGGACT CTAATCATGG	300
15	CTCTGCATGA CTCTTCCGAT TACCTGCTGG AGTCAGCCAA GATGTTTAAC TACGCGGGAT	360
	GGAAGAACAC CTGCAACAAC ATCTTCATCG TCTTCGCCAT TGTTTTATC ATCACCCGAC	420
	TGGTCATCCT GCCCTTCTGG ATCCTGCATT GCACCCTGGT GTACCCACTG GAGCTCTATC	480
20	CTGCCTTCTT TGGSTATTAC TTCCTCAATT CCATGATGGG AGTTCTACAG CTGCTGCATA	540
	TCTTCTGGGC CTACCTCAT TGTGCGATGG CCCACAAGTT CATAACTGGG AAAGCTGGTA	600
25	GAAGATGAAC GCAWGCRCGG GNAAGAAACA GAGAGCTCAG AGGGGGAGGA GGCTGCAGCT	660
	GGGGGAGGAG CAAAGAGCCG GCCCCTAGCC AATGGCCACC CCATCCTCAA TAACAACCAT	720
	CGTAAGAATG ACTGAACCAT TATTCCAGCT GCCTCCCAGA TTAATGCATA AAGCCAAGGA	780
30	ACTACCCYGC TCCCTGCGCT ATAGGGTCAC TTAAAGCTCT GGGGAAAAG GAGAAAGTGA	840
	GAGGAGAGTT CTCTGCATCC TCCCTCCTTG CTTGTACACC AGTTGCCTTT AAACCAAATT	900
35	CTAACCAGCC TATCCCCAGG TAGGGGGACG TTGGTTATAT TCTGTTAGAG GGGGACGGTC	960
	GTATTTTCCT CCCTACCCGC CAAGTCATCC TTTCTACTGC TTTTGAGGCC CTCCCTCAGC	1020
	TCTCTGTGGG TAGGGGTAC AATTCACATT CCTTATTCTG AGAATTGGC CCCAGCTGTT	1080
40	TGCCTTTGAC TCCCTGACCT CCAGAGCCAG GGTGTGCGCT TATTGTCCCA TCTGTGGGCC	1140
	TCATTCTGCC AAAGCTGGAC CAAGGCTAAC CTTTCTAAGC TCCCTAAGTT GGGCCAGAAA	1200
45	CCAAAGCTGA GCTTTTAACT TTCTCCCTCT ATGACACAAA TGAATTGAGG GTAGGAGGAG	1260
	GGTGACATA ACCCTTACCC TACCTCTGCC AAAAAGTGGG GCTGTACTG GGGACTGCTC	1320
	GGATGATCTT TCTTAGTGCT ACTTCTTTCA GCTGTGCTG TAGCGACAGG TCTAAGATCT	1380
50	GACTGCCTCC TCCTTTCTCT GGCCTCTTCC CCCTTCTCT TTTCTCTCAG CTAGGCTAGC	1440
	TGGTTTGGAG TAGAATGGCA ACTAATTCTA ATTTTATTTT ATTAAATATT TGGGGTTTGT	1500
55	GTTTTAAAGC CAGAATTACG GCTAGCACCT AGCATTTTCA CAGGAGGACC ATTTTAGACC	1560
	AAAATGTACT GTTAATGGGT TTTTTTTTAA AATTAAAAA ATTAAATAAAA AATATTAAAT	1620
60	AAAACATGGC AATAAGTGTC AGACTATTAG GAATTGAAA CAGGAGATCAA CTAATAAAC	1680

267

GAAGAG

1686

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(2) INFORMATION FOR SEQ ID NO: 123:

(i) SEQUENCE CHARACTERISTICS:

10

(A) LENGTH: 1211 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 123:

15

CAGCCTGTGC CAGACGAGGA GGTGATTGAG CTGTATGGGG GTACCCAGCA CATCCCACTA 60

TACCAGATGA GTGGCTTCTA TGGCAAGGGT CCCTCCATTA AGCAGTTCAT GGACATCTTC 120

20

TCGCTACCGG AGATGGCTCT GCTGTCTGT GTGGTGGACT ACTTCTGCGG CCACAGCCTG 180

GAGTTTGACC AACATCTCT ACAAGGACGT GACGGACGCC ATCCGAGACG TGCATGTGAA 240

25

GGGCCTCATG TACCAGTGGA TCGAGCAGGA CATGGAGAAG TACATCCTGA GAGGGGATGA 300

GACGTTTGCT GTCTGAGCC GCCTGGTGGC CCATGGGAAA CAGCTGTTCC TCATCACCAA 360

CAGTCTTTC AGCTTCGTAG ACAAGGGGAT GCGGCACATG GTGGGTCCCG ATTGGCGCCA 420

30

CTCTTCGATG TGGTCATTGT CCAGGCAGAC AAGCCCAGCT TCTTCACTGA CCGGCGCAAC 480

TTTCAGAAAA CTCGATGAGA AGGGCTCACT TCAGTGGGAC CGGATCACCC GCTTGAAAA 540

35

GGGCAAGATC TATCGGCAGG GAAACCTGTT TGACTTCTTA CGCTTGACGG AATGGCGTGG 600

CCCCCGCTG CTCTACTTCG GGGACCACCT CTATAGTGAT CTGGCGGATC TCATGCTGCG 660

GCACGGCTGG CGCACAGGCG CCATCATCCC CGAGCTGGAG CGTGAGATCC GCATCATCAA 720

40

CACGGAGCAG TACATGCACT CGCTGACGTG GCAGCAGGCG CTCACGGGCG TGCTGGAGCG 780

CATGCAGACC TATCAGGACG CGGAGTCGAG GCAGGTGCTG GCTGCCTGGA TGAAAGAGCG 840

GCAGGAGCTG AGGTGCATCA CCAAGGCCCT GTTCAATGCG CAGTTCGGCA GCATCTTCCG 900

45

CACCTTCCAC AACCCACCT ACTTCTCAAG GCGCCTCGTG CGCTTCTCTG ACCTCTACAT 960

GGCCTCCCTC AGCTGCCTGC TCAACTACCG CGTGGACTTC ACCTTCTACC CACGCCGTAC 1020

50

GCCGCTGCAG CACGAGGCAC CCCTCTGGAT GGACCAGCTT CTGCACCGGC TGATGAAGA 1080

CCCCCTCCT TGGTGACATG GCCCACATCC GCTGAGGGCA CTTTATTGT CTGGGACAGG 1140

CCCTCAGCCC CTCCTGCCCC ATCCACCCAG ACAAGCAATA AAAGTGGTCT CCTCCCTGAA 1200

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AAAAAAAAA A 1211

60

(2) INFORMATION FOR SEQ ID NO: 124:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 1804 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 124:

10 CGCACCTATG GGCTCGCTAC CAGGACATGC GGAGACTGGT GCACGACCTC CTGCCCCCGG 60
 AGGTCTGCAG TCTCCTGAAC CCAGCAGCCA TCTACGCCAA CAACGAGATC AGCCTGCGTG 120
 15 ACGTTGAGGT CTACGGCTTT GACTACGACT ACACCCTGGC CCAGTATGCA GACGCACTGC 180
 ACCCCGAGAT CTTCACTACC GCCCGTGACA TCCTGATCGA GCACTACAAG TACCCAGAAG 240
 GGATTCCGAA GTATGACTAC AACCCAGCT TTGCCATCCG TGGCCTCCAC TATGACATTC 300
 20 AGAAGAGCCT TCTGATGAAG ATTGACGCCT TCCACTACGT GCAGCTGGGG ACAGCCTACA 360
 GGGGCCTCCA GCCTGTGCCA GACGAGGAGG TGATTGAGCT GTATGGGGGT ACCCAGCACA 420
 25 TCCCACTATA CCAGATGAGT GGCTTCTATG GCAAGGGTCC CTCCATTAAG CAGTTCATGG 480
 ACATCTTCTC GCTACCGGAG ATGGCTCTGC TGTCTGTGT GGTGGACTAC TTTCTGGGCC 540
 ACAGCCTGGN AGTTTGACCA AGCACATCTC TACAAGGACG TGACGGACGC CATCCGAGAC 600
 30 GTGCATGTGA AGGGCCTCAT GTACCACTGG ATCGAGCAGG ACATGGAGAA GTACATCCTG 660
 AGAGGGGATG AGACGTTTGC TGTCTGAGC CGCCTGGTGG CCCATGGGAA ACAGCTGTTC 720
 35 CTCATCACCA ACAGTCCTTT CAGCTTCGTA GACAAGGGGA TGCGGCACAT GGTGGGTCCC 780
 GATTGGCGCC ACTCTTCGAT GTGGTCATTG TCCAGGCAGA CAAGCCCAGC TTCTTCACTG 840
 ACCGGCGCAA GCTTTTCAGA AAACTCGATG AGAAGGGCTC ACTTCAGTGG GACCGGATCA 900
 40 CCCGCTTGA AAAGGGCAAG ATCTATCGGC AGGGAACCT GTTTGACTTC TTACGCTTGA 960
 CGGAATGGCG TGGCCCCCGC GTGCTCTACT TCGGGACCA CCTCTATAGT GATCTGGCGG 1020
 45 ATCTCATGCT GCGGCACGGC TGGCGCACAG GCGCCATCAT CCCCAGAGCTG GAGCGTGAGA 1080
 TCCGCATCAT CAACACGGAG CAGTACATGC ACTCGCTGAC GTGGCAGCAG GCGCTCACGG 1140
 GGCTGCTGGA GCGCATGCAG ACCTATCAGG ACGCGGAGTC GAGGCAGGTG CTGGCTGCCT 1200
 50 GGATGAAAGA GCGGCAGGAG CTGAGGTGCA TCACCAAGGC CCTGTTCATG GCGCAGTTCTG 1260
 GCAGCATCTT CCGCACCTTC CACAACCCCA CCTACTTCTC AAAGGCGCCT CGTGGCCTTC 1320
 55 TCTGACCTCT ACATGGCCTC CCTCAGCTGC CTGCTCAACT ACCGCGTGGG CTTCACCTTC 1380
 TACCCACGCC GTACGCCGCT GCAGCACGAG GCACCCCTCT GGATGGACCA GCTCTGCACC 1440
 GGCTGCATGA AGACCCCTT CCTTGGTGAC ATGGCCGACA TCCGCTGAGG GCACCTTTAT 1500
 60

TGTCTGGGAC AGGCCCTCAG CCCCTCCTGC CCCATCCACC CAGACAAGCA ATAAAAGTGG 1560
 TCTCCTCCCT GTGCATGCTT CTGCTTTCAG CCCAGCCTC GTCATTGAC TGTGAGGATC 1620
 5 CTCTGGGTGT CAGGGAAGTC CTCCTCCAGC AGTGAGTCAT CGAAGGGTTC AAAAAAGGTG 1680
 TCGCTGCCAA AGACAGGGTT GGGGACAGAG ACCAGGGTGG GGTGGTCCC TTCTTGCCAC 1740
 10 GGTGAGAAGT CGTCGTCAGC CGGACGCGTG GGTGACCCG GGAATTCCGG ACCGGTACCT 1800
 GCAG 1804

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(2) INFORMATION FOR SEQ ID NO: 125:

(i) SEQUENCE CHARACTERISTICS:

20

- (A) LENGTH: 1282 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 125:

25

CCGCAGGNCA GCGACGCGAC TCTGGTGCGG GCCGTCTTCT TCCCCCGAG CTGGGCGTGC 60
 GCGGCCGCAA TGAAGTGGGA GCTGCTGCTG TGGCTGCTGG TGCTGTGCGC GCTGCTCCTG 120
 30 CTCTTGGTGC AGCTGCTGCG CTTCTGAGG GCTGACGGCG ACCTGACGCT ACTATGGGCC 180
 GAGTGGCAGG GACGACGCCC AGAATGGGAG CTGACTGATA TGGTGGTGTG GGTGACTGGA 240
 35 GCCTCGAGTG GAATTGGTGA GGAGCTGGCT TACCAGTTGT CTAAACTAGG AGTTTCTCTT 300
 GTGCTGTCAG CCAGAAGAGT GCATGAGCTG GAAAGGGTGA AAAGAAGATG CCTAGAGAAT 360
 GGCAATTTAA AAGAAAAAGA TATACTTGTT TTGCCCCCTG ACCTGACCGA CACTGGTTCC 420
 40 CATGAAGCGG CTACCAAAGC TGTTCCTCAG GAGTTTGGTA GAATCGACAT TCTGGTCAAC 480
 AATGGTGGAA TGCCCAGCG TTCTCTGTGC ATGGATACCA GCTTGGATGT CTACAGAAAG 540
 CTAATAGAGC TTAAGTACTT AGGGACGGTG TCCTTGACAA AATGTGTTCT GCCTCACATG 600
 45 ATCGAGAGGA AGCAAGGAAA GATTGTTACT GTGAATAGCA TCCTGGGTAT CATATCTGTA 660
 CCTCTTTCCA TTGGATACTG TGCTAGCAAG CATGCTCTCC GGGGTTTTTT TAATGGCCTT 720
 50 CGAACAGAAC TTGCCACATA CCCAGGTATA ATAGTTTCTA ACATTTGCCC AGGACCTGTG 780
 CAATCAAATA TTGTGGAGAA TTCCCTAGCT GGAGAAGTCA CAAAGACTAT AGGCAATAAT 840
 GGAGACCACT CCCACAAGAT GACAACCACT CGTTGTGTGC GGCTGATGTT AATCAGCATG 900
 55 GCCAATGATT TGAAAGAAGT TTGGATCTCA GAACAACCTT TCTTGTTACT AACATATTTG 960
 TGGCAATACA TGCCAACCTG GGCCTGGTGG ATAACCAACA AGATGGGGA GAAAAGGATT 1020
 60 GAGAACTTTA AGAGTGGTGT GGATGCAGAC TCTTCTTATT TTAAAATCTT TAAGACAAAA 1080

CATGACTGAA AAGAGCAYCT GTACTTTTCA AGCCACTGGA GGGARAAATG GAAAACATGA 1140
 AAACAGCAAT CTTCTTATGC TTCTGAATAA TCAAAGACTA ATTTGTGRTT TTA CTTTCTTTTA 1200
 5 ATAGATATGA CTTTGCTTCC AACATGGAAT GAAATAAAAA ATAAATAATA AAAGATTGCC 1260
 ATGGAAAAAA AAAAGNNGGG AN 1282

10

(2) INFORMATION FOR SEQ ID NO: 126:

15

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 1296 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 126:

GGCAGAGCTT AGAGTGTTGA AAAGGCAACC AGGTTGGCCG TAAGTGCCTG CTGGAATGCG 60
 25 TGTGCCTCCA CASGGRTCTG GGCATCCGGA CTGATAACCA GCCGGCCAGA CTGAGGGATG 120
 GAAGGCACTG AGATGGGGGC CCGTCCAGGC GGACACCCGC AGAAATGGAG CTTTCTGTGG 180
 TCTCTTGAC TCTGGCTGCC TCTTGCCCTC TCTGTGTCTC TCTTCTTTGG TCTCTCCCTC 240
 30 TCTCTCTCTC AGCCTGGTCT TTCTCTTTGG TGCACACTTA GTTATTGTGTG TGAGCAATGG 300
 AAGTTCAAAG GAACTCCCTC TCCAGCTCTT CTGAATCTTG GGACACAGCC TAAAAAGGAC 360
 35 AAAAAAGTTAG AAGACAGCAT AGCAACTCAG CTCAGGGRGC TACCAGAGAA AAATAGCAAC 420
 TGATGTGGGT GCTTTTTTTT TTTTTTAAT TTGAATAAAA AGAATTAGAA GTGATGTCCT 480
 TTTATAAAAT GCCTTCTCCC CCTTCCCGCC TACAGTCTCT TCCTCTCCCC TTAGAGGGGG 540
 40 GAAAGTGTAT AAACCTACAG GGTGTGAGT CTGAAAAGAG GATCCCCCTC ACCCCCACCC 600
 TGGGCAGAGC AGTGGGGGTT GGGGGGTGGG AGAGGGGGAC ACAGATCCTG GCACACTGTG 660
 45 GATATTTCTT GCAGATTGCA GTCTCTTGTG GCCCAAACAG GTTAGGTAGA CTATCGCCTC 720
 TGGCAGGTGC CACCTTTTGG TACCAACATG TTCTGAGGTG TTAGGATTTG GGTGGGTTT 780
 TTTTGTTTT TTTTTTTTTT CCNTTTGGTC TTTTTTTTTT TCYCCTTKTA AAGAAAAGCT 840
 50 AAAGGCCGCT GTGAGTCTG GTGGCAGGCT CTCCATGGAT GTAGCATATC GAAGATAATT 900
 TTTATACTGC ATTTTATATG ATTATTTTGT AATGTGTGAT TCCGTCTGCT GAGGAGGTGG 960
 55 GAGGGGCTCC AGGGAAGCC ACCCACCTTC AGTGAGGTTG CTCCCAGCT GAGCGCACCG 1020
 GGCATGGGAT GTGGAGGCTG GCGACACACC CTGTGCTCTT CCAAGGCTGG GCGCGTGGG 1080
 CGTCCAGAGT CTCTCTGGGT CTCAGATGTC CATCTGCCAC CTCTTGTTAA GGCTCTAGCC 1140
 60

271

AGAAGGGAGG GTGAGGGTAG AAGAAAGTTA TTCCCGAAGA AAAAAAGAAT GAAAAGTCAT 1200
 TGTACTGAAC TGTTTTATA TTTTAAAAG TTAATTTTA AAGCGGACGT CGTGGGTGCA 1260
 5 CCCGGAATT CCCGGACCGG TACTGTCAGG TCTAAC 1296

10 (2) INFORMATION FOR SEQ ID NO: 127:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 737 base pairs
 (B) TYPE: nucleic acid
 15 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 127:

20 GGCANAGTGG AGGCAATGCC AGCTCCAGGA CAGAGGCTCA GGTGCCCAAC GGGCAAGGCA 60
 GCCCAGGGGG CTGTGTCTGT TCAAGTCAGG CTCCCCGGC CCYTCGCGCA NCAGCGCTTC 120
 CACGGGCAGC CCGGGGCCCC ACCCCACGCA CTGAAGAGGC CGCCTGGGCT GCCATGGCCC 180
 25 TGACCTTCCT GCTGGTGCTG CTCACCCTGG CCACGCTCTG CACACGGCTG CACAGAACT 240
 TCCGACGCGG GGAGAGCATC TACTGGGGGC CCACAGCGGA CAGCCAGGAC ACACTGGCTG 300
 30 CTGTGCTGAA GCGGAGGCTG CTGCAGCCCT CGCGCCGGGT CAAGCGCTCG CGCCGGAGAC 360
 CCYTCYTCCC GCCCAGCCG GACAGCGGCC CGGAAGGCGA GAGCTCGGAG TGACGGCCTG 420
 GGACCTGCCA CTGTGGCGTG CGGTCTCCCC GCGCCGCGAG GCCGCGAMCT NTGCCACGTG 480
 35 GACCGCGCGC NGGGCGCTMC CCTGGTGGCG ATGGCGCGG ACTGGCGAGC ACTGCGKGG 540
 CTTTCCTCCT TGTGGTTGC TGAGTGGGCG GCCAAGGGGA GAAAAGGAGC CGCTTYTGCC 600
 40 TCCCTTGCCA AAACCTCCGT TCTAATTAAA TTATTTTATAG TAGAAAAAAA AAAAAAAA 660
 AAAAAAAA AAAAAAAA AAAAAAAC TCGAGGGGGG GCGCGGTACC CAATTNGCCA 720
 AATAGCGATC GTATNAA 737
 45

50 (2) INFORMATION FOR SEQ ID NO: 128:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1925 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 128:

CCCCGCTCC AAAGCTAACC CTCGGGCTTG AGGGGAAGAR CCTGACTGTA CGTTCCTTCT 60
 60

[illegible]

ACTGCGGCCG CTGTCCCTTC TGTCGTCTTC TCGCAGCCGT ACCCTTCTGT CGTCTTCTCG 1920
CAGCC 1925

5

(2) INFORMATION FOR SEQ ID NO: 129:

10

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2713 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 129:

TCCTACCTTC CCAACCCTCT GGCATCCCCA GCACTGATGG TCCTGGCATC CACGGCTGAG 60
20 GCCAGCCGTG ACTGCTTCCA TCCCTTGTC A GACCCACGA CCCTTTGGTG TACCTGTYTC 120
AGTTGACAAG GACGTGCATA TTCCTTTCAC CAACGGTTCC TATACCTTTG CCTCTATGTA 180
25 CCATCGGCAA GGTGGGGTGC CAGGCACTTT TGCCAATCGT GATTTCCCCC CTTCTCTACT 240
ACACCTCCAC CCTCAATTG CTCCCCAAA TCTAGATTGC ACCCAATCA GTATGCTGAA 300
TCATAAGTGG TGTGGGGT TCCGGCCTTT GSCTCCACCC GRGGACCGGG RGAGYTATCA 360
30 GTCAGCTTTA CGCCGGCCAA GCGACTTAAG AACTGCCATG ACACAGAGTC TCCCCACTTG 420
CGCNTCTCAG ATGCAGATGG GAANGAATAT GACTTTGGGA CACAGCTGCM ATCTAGCTCC 480
35 CCCGGTTCAC TAAAGGTTGA TGACACTGGG AAGAAGATTT TTGCTGTCTC TGGCCTCAT 540
TCTGATCGGG AAGCCTCATC TAGCCCAGAG GNTCGGNAAT GACAGATGTA AGAAGAAAGC 600
AGCGGCATTG TTCGACAGCC AGGCCCCAAT TTGCCCCATC TGCCAGGTCC TGCTGAGGCC 660
40 CAGTGAGCTG CAGGAGCATA TGGAGCAGGA ACTGGAGCAG CTAGCCCAAC TGCCCTCGAG 720
CAAGAATTCC CTTCTGAAGG ATGCCATGGC TCCAGGCACC CCAAAGTCCC TCCTGTGTGTC 780
TGCTTCCATC AAGAGGGAAG GAGAGTCTCC AACGGCATCA CCCCCTCAT CTGCCACCGA 840
45 TGACCTCCAC CATTCAGACA GATACCAGAC CTTTCTCCGA GTACGAGCCA ACCGGCAGAC 900
CCGAYTGAAT GYTCGGATTG GGAAAATGAA ACGGAGGAAG CAAGATGAAG GGCAGGTATG 960
50 TCCCCGTGTC AACCGCCCCC TGGCAGGATC GGAGCAGGAG ATGAGTAGGC ATGTGAGCA 1020
TTGCCTTTCT AAGAGGGAAG GCTCCTGCAT GGCTGAGGAT GATGCTGTGG ACATCGAGCA 1080
TGAGAACAAC AACCGCTTTG AGGAGTATGA GTGGTGTGGA CAGAAGCTGA TACGGGCCAC 1140
55 CACTCTCCTG GAAGGTGGCT TCCGAGGCTC TGCTTCTGAT ATGTGAGGCG GCAAAGAGAA 1200
CCCGACAGT GATGCTGACT TGGATGTGGA TGGCGATGAC ACTCTGGAGT ATGGAAGGCC 1260
60 ACAATACACA GAGGCTGATG TCATCCCCTG CACAGGCGAG GAGCCTGGTG AAGCCAAGGA 1320

274

	GAGAGAGGCA CTTCGGGGCG CAGTCCTAAA TGGCGGCCCT CCCAGCACGC GCATCACACC	1380
5	TGAGTTCTCT AAATGGGCCA GTGATGAGAT GCCATCCACC AGCAATGGTG AAAGCAGCAA	1440
	GCAGGAGGCC ATGCAGAAGA CCTGCAAGAA CAGCGACATC GAGAAAATCA CCGAAGATTC	1500
	AGCTGTGACC ACGTTTGAGG CTCTGAAGGC TCGGGTCAGA GAACTTGAAC GGCAGCTATC	1560
10	TCGTGGGGAC CGTTACAAAT GCCTCATCTG CATGGACTCG TACTCGATGC CCCTAACGTC	1620
	CATCCAGTGT TGGCACGTGC ACTGCGAGGA GTGCTGGCTG CGGACCCTGG GTGCCAAGAA	1680
15	GCTCTGCCCT CAGTGCAACA CGATCACAGC GCGCGGAGAC CTGCGGAGGA TCTACTTGTC	1740
	AGCTATCTGC CCCAGGCAGG CCTCGCCTCC AGCAGCCCCA CCTGCCCCCA GCCTCTGTGA	1800
	CAGTGACCGT YTCCCTTTGT ACATACTTGC ACACAGGTTT CCCATGTACA TACATGCACA	1860
20	TACTCAAACA TGCGTACACA CACACACATT TACACACGCA GGACTCTGGA GCCAGAGTAG	1920
	AGGCTGTGGC CCAGGCACTA CCTGCTGGCT CCCACCTATG GTTTGGGGGC CATACCTGTT	1980
25	CCAGCTCTGT TCCCAGGGTG GGGCAGGGAG GTGGGGTTG GGGGAGTAGT GGGGCACGGC	2040
	TCCTAAGATC CAGCCCCCAT ACTGACAGAC GGACAGACAG ACATGCAAAC ACCAGACTGA	2100
	AGCACATGTA ATATAGACCG TGTATGTTTA CAATGTTGTG TATAAATGGG ACAACTCCTC	2160
30	GCCCTCTACC TGTCCCCTCC CCCTTTGGTT GTATGATTTT CTTCTTTTTT AAGAACCCCT	2220
	GGAAGCAGCG CCTCCTTCAG GGTGCGCTGG GAGCTCGGCC CATCCACCTC TTGGGGTAYC	2280
35	TGCCTCTCTC TCTCCTGTGG TGTCCCTTCC CTCTCCCATG TGCTCGGTGT TCAGTGGTGT	2340
	ATATTCTTTC TCCCAGACAT GGGGCACACG CCCCAGGGA CATGATCCTC TCCTTAGTCT	2400
	TAGCTCATGG GGCTCTTTAT AAGGAGTTGG GGGGTAGAGG CAGGAAATGG GAACCGAGCT	2460
40	GAAGCAGAGG CTGAGTTAGG GGGCTAGAGG ACAGTGCTCC TGGCCACCCA GCCTCTGCTG	2520
	AGAACCATTG CTGGGATTAG AGCTGCCTTT CCCAGGGAAA AAGTGTCTGC TCCCCGACCC	2580
45	TCCCGTGGG CCTGTGGTGT GATGCTGTGT CTGTATATTC TATACAAAGG TACTTGTCCT	2640
	TTCCCTTTGT AACTACATT TGACATGGAT TAAACCAGTA TAAACAGTTA AAAAAAAAAA	2700
	AAAAAAAACT CGA	2713
50		

(2) INFORMATION FOR SEQ ID NO: 130:

- 55 (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 1011 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear
- 60

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 130:

	AGAGGACGGT GTGACCCGGG AGGAAGTAGA GCCTGAGGAG GCTGAAGAAG GCATCTCTGA	60
5	GCAACCCTGC CCAGCTGACA CAGAGGTGGT GGAAGACTCC TTGAGGCAGC GTAAAAGTCA	120
	GCATGCTGAC AAGGGACTGT AGATTTAATG ATGCGTTTTC AAGAATACAC ACCAAAACAA	180
10	TATGTCAGCT TCCCTTTGGC CTGCAGTTTG TACCAAATCC TTAATTTTTY YTGAATGAGC	240
	AAGCTTCTCT TAAAAGATGC TCTCTAGTCA TTTGGTCTCA TGGCAGTAAG CCTCATGTAT	300
	ACTAAGGAGA GTCTTCCAGG TGTGACAATC AGGATATAGA AAAACAAACG TAGTGTNTGG	360
15	GATCTGTTTG GAGACTGGGA TGGGAACAAG TTCATTTACT TAGGGGTCAG AGAGTCTCGA	420
	CCAGAGGAGG CCATTCCCAG TCCTAATCAG CACCTTCCAG AGACAAGGCT GCAGGCCCTG	480
20	TGAAATGAAA GCCAAGCAGG AGCCTTGGCT CTGAGNCATC CCCAAAGTGT AACGTAGAAG	540
	CCTTGCATCC TTTTCTTGTG TAAAGTATTT ATTTTGTGCA AATTGCAGGA AACATCAGGC	600
	ACCACAGTGC ATGAAAAATC TTTCACAGCT AGAAATTGAA AGGGCCTTGG GTATAGAGAG	660
25	CAGCTCAGAA GTCATCCCAG CCCTCTGAAT CTCCTGTGCT ATGTTTATT TCTTACCTTT	720
	AATTTTCCA GCATTTCCAC CATGGGCATT CAGGCTCTCC AACTCTTCA CTATTATCTC	780
30	TTGGTCAGAG GACTCCAATA ACAGCCAGGT TTACATGAAC TGTGTTTGTG CATCTGACC	840
	TAAGGGGTTT AGATAATCAG TAACCATAAC CCCTGAAGCT GTGACTGCCA AACATCTCAA	900
	ATGAAATGTT GTRGCCATCA GAGACTCAAA AGGAAGTAAG GATTTTACAA GACAGATTAA	960
35	AAAAAATTG TTTGTGCCAA AAAANAAAAA AAAAAACTC GAAGGGGGGG C	1011

40 (2) INFORMATION FOR SEQ ID NO: 131:

(i) SEQUENCE CHARACTERISTICS:

- | | |
|----|-----------------------------|
| | (A) LENGTH: 2278 base pairs |
| | (B) TYPE: nucleic acid |
| 45 | (C) STRANDEDNESS: double |
| | (D) TOPOLOGY: linear |

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 131:

50	GTAATTCGGC ACGAGGCGCC CAACATGGCG GGTGGGCGTT ATGGCCCGCA SCTAACGGCG	60
	CTCCTGGCCG CCTGGATCGC GGCTGTGGCG GCGACGAAAT GAGGCGAGGA GGCCGCGCTG	120
	CCGCCGGAGC AGAGCCGGGT CCAGCCCATG ACCGCCCTTA ACCTGAGGTT GGTGATGGAG	180
55	GGCGAGTGGA TGCTGAAATT TTACGCCCCA TGGTGTCTTT TTTTACGAGCA GACTGATTCA	240
	GAATGGGAGG CTTTTCGAAA GAATGGTGAA ATACTTAAAT TTTTGTCTGG GAAGGTAGAT	300
60	GTCATTCAAG AACCAGGTTT GAGTGGCCGC TTCTTTTCT AACTTCTCTGC AGCATTTTTT	360

	CATGCAAAGG ATGGGATATT CCGCCGTTAT CGTGGCCCAG GAATCTTCGA AGACCTGCAG	420
5	AATTATATCT TAGAGAAGAA ATGGCAATCA GTCGAGCCTC TGA CTGGCTG GAAATCCCCG	480
	GCTTCTCTAA CGATGTCTGG AATGGCTGGT CTTTTTAGCA TCTCTGGCAA GATATGGCAT	540
	CTTCACA ACT ATTTACAGT GACTCTTGGA ATTCTGCTT GGTGTTCTTA TGTCTTTTTC	600
10	GTCATAGCCA CCTTGGTTTT TGGCCTTTTT ATGGGTCTGG TCTTGGTGGT AATATCAGAA	660
	TGTTTCTATG TGCCACTTCC AAGGCATTTA TCTGAGCGTT CTGAGCAGAA TCGGAGATCA	720
15	GAGGAGGCTC ATAGAGCTGA ACAGTTGCAG GATGCGGAGG AGGAAAAAGA TGATTCAAAT	780
	GAAGAAGAAA ACAAAGACAG CCTTGTAGAT GATGAAGAAG AGAAAGAAGA TCTTGGCGAT	840
	GAGGATGAAG CAGAGGAAGA AGAGGAGGAG GACAAC TTGG CTGCTGGTGT GGATGAGGAG	900
20	AGAAGTGAGG CCAATGATCA GGGCCCCCA GGAGAGGACG GTGTGACCCG GGAGGNAAGT	960
	AGAGCCTGAG GAGGCTGAAG AAGGCATCTC TGAGCAACCC TGCCAGCTG ACACAGAGGT	1020
25	GGTGAAGAC TCCTTGAGGC AGCGTAAAAG TCAGCATGCT GNCAAGGGAC TGTAGATTTA	1080
	ATGATGCGTT TTCAAGAATA CACACCAAAA CAATATGTCA GCTTCCCTTT GGCCTGCAGT	1140
	TTGTACCAAA TCCTTAATTT TTCTGAATG AGCAAGCTTC TCTTAAAAGA TGCTCTCTAG	1200
30	TCATTTGGTC TCATGGCAGT AAGCCTCATG TATACTAAGG AGAGTCTTCC AGGTGTGACA	1260
	ATCAGGATAT AGAAAAACAA ACGTAGTGTN TGGGATCTGT TTGGAGACTG GGATGGGAAC	1320
35	AAGTTCATTT ACTTAGGGGT CAGAGAGTCT CGACCAGAGG AGGCCATTCC CAGTCCTAAT	1380
	CAGCACCTTC CAGAGACAAG GCTGCAGGCC TGTGAAATGA AAGCCAAGCA GGAGCCTTGG	1440
	CTCTGAGGCA TCCCCAAAGT GTAACGTAGA AGCCTTGCA TCTTTTCTTG TGTAAGTAT	1500
40	TTATTTTGT CAAATTGCAG GAAACATCAG GCACCACAGT GCATGAAAAA TCTTTCACAG	1560
	CTAGAAATTG AAAGGGCCTT GGGTATAGAG AGCAGCTCAG AAGTCATCCC AGCCCTCTGA	1620
45	ATCTCCTGTG CTATGTTTTA TTTCTTACCT TTAATTTTTC CAGCATTTCC ACCATGGGCA	1680
	TTCAGGCTCT CCACACTCTT CACTATTATC TCTTGGTACG AGGACTCCAA TAACAGCCAG	1740
	GTTTACATGA ACTGTGTTTG TTCATTCTGA CCTAAGGCTT TTAGATAATC AGTAACCATA	1800
50	ACCCCTGAAG CTGTGACTGC CAAACATCTC AAATGAAGAT TTGTRGCCAT CAGAGACTCA	1860
	AAAGGAAGTA AGGATTTTAC AAGACAGATT AAAAAAAT TTTTCTCTCC NAAAAATAG	1920
55	TTGTGTTGA TTTTTTTTTA AGTTTCTAA GCAATATTTT TAAAGGAGGA AGTCCTCTAA	1980
	GTCTTGCCAG TACAAGGTAG TCTTGTGAAG AAAATTTTAA TATCTTTTTG TTTTCATCTC	2040
	AAGGGGTTCC CTGGGTCTTG AACTACTTTA ATAATAATA AAAAAAGCACT TCTGATTTTC	2100
60	CTTCAGTGAT GTGCTTTTGG TGAAAGAATT AATGAAGCTT AATACCTGAA AGTGAAAGAT	2160

	TTGATTTTGT TTCCATCTTC TGTAATCTTC CAAAGAATTA TATCTTTGTA AATCTCTCAA	2220
5	TACTCAATCT ACTGTAAGTA CCCAGGGRGG STAATTTCYT TAAAAAAAAA AAAAAAAA	2278
(2) INFORMATION FOR SEQ ID NO: 132:		
10	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1088 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
15	(D) TOPOLOGY: linear	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 132:		
20	GGCAGGGGCG GCGTGAACCC GTCGGGCACT GTGTCCCTGA CAATGGGAAC AGCCGACAGT	60
	GATGAGATGG CCCCGGAGCC CCACAGCACA CCCACATCGA TGTGCACATC CACCAGGAGT	120
	CTGCCCTGGC CAAGCTCCTG CTCACCTGCT GCTCTGCGCT GCGGCCCCGG GCCACCCAGG	180
25	CCAGGGGCAG CANCCGGCTG CTGGTGGCCT CGTGGGTGAT GCAGATCGTG CTGGGGATCT	240
	TGAGTGCAGT CCTAGGAGGA TTTTCTACA TCCGCGACTA CACCCTCCTC GTCACCTCGG	300
	GAGCTGCCAT CTGGACAGGG GCTGTGGCTG TGCTGGCTGG AGCTGCTGCC TTCATTTAYG	360
30	AGAAACGGGG TGGTACATAC TGGGCCCTGC TGAGGACTCT GCTARCGCTG GCAGCTTTCT	420
	CCACAGCCAT CGCTGCCCTC AAACCTTGGG ATGAAGATTT CCGATATGGC TACTCTTATT	480
35	ACAACAGTGC CTGCCGCATC TCCAGCTCGA GTGACTGGAA CACTCCAGCC CCCACTCAGA	540
	GTCCAGAAGA AGTCAGAAGG CTACACCTAT GTACCTCCTT CATGGACATG CTGAAGGCCT	600
	TGTTCAGAAC CCTTCAGGCC ATGCTCTTGG GTGTCTGGAT TCTGCTGCTT CTGGCATCTC	660
40	TGGCCCTCT GTGGCTGTAC TGCTGGAGAA TGTTCCTAAC CAAAGGGAAA AGAGACCAGA	720
	AGGAAATGTT GGAAGTGAGT GGAATCTAGC CATGCCTCTC CTGATTATTA GTGCCTGGTG	780
45	CTTCTGCACC GGGCGTCCCT GCATCTGACT GCTGGAAGAA GAACCAGACT GAGGAAAAGA	840
	GGCTCTTCAA CAGCCCCAGT TATCCTGGCC CCATGACCGT GGCCACAGCC CTGCTCCAGC	900
	AGCACTTGCC CATTCTTAC ACCCCTTCCC CATCCTGCTC CGCTTCATGT CCCCTCCTGA	960
50	GTAGTCATGT GATAATAAAC TCTCATGTTA TTGTTCCNAA AAAAAAAAAA AAAAAAAAT	1020
	TGGGGGGGGG CCGGTACCCA TTGGGCCTNN GGGGGNGGTT TAAAAATTAAT GGGGGGGGTT	1080
55	TAAAAGGG	1088

60 (2) INFORMATION FOR SEQ ID NO: 133:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 553 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 133:

10 GGCAGAGAGC AGATGGCCTT GACACCAGCA GGGTGACATC CGCTATTGCT ACTTCTCTGC 60
TCCCCACAG TTCCTCTGGA CTTCTCTGGA CCACAGTCCT CTGCCAGACC CCTGCCAGAC 120
15 CCCAGTCCAC CATGATCCAT CTGGGTCACA TCCTCTTCCT GCTTTTGCTC CCAGTGGCTG 180
CAGCTCAGAC GACTCCAGGA GAGAGATCAT CACTCCCTGC CTTTACCCT GGCACCTCAG 240
GCTCTTGTTT CGGATGTGGG TCCCTCTCTC TGCCGCTCCT GGCAGGCCTC GTGGCTGCTG 300
20 ATGCGGTGGC ATCGCTGCTC ATCGTGGGGG CGGTGTTCTT GTGCGCACGC CCACGCCGCA 360
GCCCCGCCCA AGATGGCAAA GTCTACATCA ACATGCCAGG CAGGGGCTGA CCCTCCTGCA 420
GCTTGGACCT TTGACTTCTG ACCCTCTCAT CCTGGATGGT GTGTGGTGGC ACAGGAACCC 480
25 CCGCCCCAAC TTTTGGATTG TAATAAAACA ATTGAAACAC CAAAAAAAAA AAAAAAAAAA 540
AAAAAAAAAA AAA 553

30

(2) INFORMATION FOR SEQ ID NO: 134:

35 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 467 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 134:

40 Met Arg Pro Gln Glu Leu Pro Arg Leu Ala Phe Pro Leu Leu Leu Leu
1 5 10 15
Leu Leu Leu Leu Leu Pro Pro Pro Pro Cys Pro Ala His Ser Ala Thr
45 20 25 30
Arg Phe Asp Pro Thr Trp Glu Ser Leu Asp Ala Arg Gln Leu Pro Ala
35 40 45
50 Trp Phe Asp Gln Ala Lys Phe Gly Ile Phe Ile His Trp Gly Val Phe
50 55 60
Ser Val Pro Ser Phe Gly Ser Glu Trp Phe Trp Trp Tyr Trp Gln Lys
55 65 70 75 80
Glu Lys Ile Pro Lys Tyr Val Glu Phe Met Lys Asp Asn Tyr Pro Pro
85 90 95
60 Xaa Phe Lys Tyr Glu Asp Phe Gly Pro Leu Phe Thr Ala Lys Phe Phe
100 105 110

279

	Asn	Ala	Asn	Gln	Trp	Ala	Xaa	Ile	Phe	Gln	Ala	Ser	Gly	Ala	Lys	Tyr	
			115					120					125				
5	Ile	Val	Leu	Thr	Ser	Lys	His	His	Glu	Gly	Phe	Thr	Leu	Trp	Gly	Ser	
		130					135					140					
	Glu	Tyr	Ser	Trp	Asn	Trp	Asn	Ala	Ile	Asp	Glu	Gly	Pro	Lys	Arg	Asp	
	145					150					155					160	
10	Ile	Val	Lys	Glu	Leu	Glu	Val	Ala	Ile	Arg	Asn	Arg	Thr	Asp	Leu	Arg	
				165						170					175		
	Phe	Gly	Leu	Tyr	Tyr	Ser	Leu	Phe	Glu	Trp	Phe	His	Pro	Leu	Phe	Leu	
15				180						185				190			
	Glu	Asp	Glu	Ser	Ser	Ser	Phe	His	Lys	Arg	Gln	Phe	Pro	Val	Ser	Lys	
		195						200					205				
20	Thr	Leu	Pro	Glu	Leu	Tyr	Glu	Leu	Val	Asn	Asn	Tyr	Gln	Pro	Glu	Val	
		210					215					220					
	Leu	Trp	Ser	Asp	Gly	Asp	Gly	Gly	Ala	Pro	Asp	Gln	Tyr	Trp	Asn	Xaa	
	225					230					235					240	
25	Thr	Gly	Phe	Leu	Ala	Trp	Leu	Tyr	Asn	Glu	Ser	Pro	Val	Arg	Gly	Thr	
				245						250					255		
	Val	Val	Thr	Asn	Asp	Arg	Trp	Gly	Ala	Gly	Ser	Ile	Cys	Lys	His	Gly	
30				260					265					270			
	Gly	Phe	Tyr	Thr	Cys	Ser	Asp	Arg	Tyr	Asn	Pro	Gly	His	Leu	Leu	Pro	
			275					280					285				
35	His	Lys	Trp	Glu	Asn	Cys	Met	Thr	Ile	Asp	Lys	Leu	Ser	Trp	Gly	Tyr	
		290					295					300					
	Arg	Arg	Glu	Ala	Gly	Ile	Ser	Asp	Tyr	Leu	Thr	Ile	Glu	Glu	Leu	Val	
	305					310					315					320	
40	Lys	Gln	Leu	Val	Glu	Thr	Val	Ser	Cys	Gly	Gly	Asn	Leu	Leu	Met	Asn	
				325						330					335		
	Ile	Gly	Pro	Thr	Leu	Asp	Gly	Thr	Ile	Ser	Val	Val	Phe	Glu	Glu	Arg	
45				340					345					350			
	Leu	Arg	Gln	Met	Gly	Ser	Trp	Leu	Lys	Val	Asn	Gly	Glu	Ala	Ile	Tyr	
		355						360					365				
50	Glu	Thr	His	Thr	Trp	Arg	Ser	Gln	Asn	Asp	Thr	Val	Thr	Pro	Asp	Val	
		370					375					380					
	Trp	Tyr	Thr	Ser	Lys	Pro	Lys	Glu	Lys	Leu	Val	Tyr	Ala	Ile	Phe	Leu	
	385					390					395					400	
55	Lys	Trp	Pro	Thr	Ser	Gly	Gln	Leu	Phe	Leu	Gly	His	Pro	Lys	Ala	Ile	
				405						410				415			
	Leu	Gly	Ala	Thr	Glu	Val	Lys	Leu	Leu	Gly	His	Gly	Gln	Pro	Leu	Asn	
60				420					425					430			

280

Trp Ile Ser Leu Glu Gln Asn Gly Ile Met Val Glu Leu Pro Gln Leu
 435 440 445

5 Thr Ile His Gln Met Pro Cys Lys Trp Gly Trp Ala Leu Ala Leu Thr
 450 455 460

Asn Val Ile
 465

10

(2) INFORMATION FOR SEQ ID NO: 135:

15 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 222 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 135:

Met Trp Ser Ala Gly Arg Gly Gly Ala Ala Trp Pro Val Leu Leu Gly
 1 5 10 15

25 Leu Leu Leu Ala Leu Leu Val Pro Gly Gly Gly Ala Ala Lys Thr Gly
 20 25 30

Ala Glu Leu Val Thr Cys Gly Ser Val Leu Lys Leu Leu Asn Thr His
 35 40 45

30 His Arg Val Arg Leu His Ser His Asp Ile Lys Tyr Gly Ser Gly Ser
 50 55 60

Gly Gln Gln Ser Val Thr Gly Val Glu Ala Ser Asp Asp Ala Asn Ser
 65 70 75 80

35 Tyr Trp Arg Ile Arg Gly Gly Ser Glu Gly Gly Cys Arg Arg Gly Ser
 85 90 95

40 Pro Val Arg Cys Gly Gln Ala Val Arg Leu Thr His Val Leu Thr Gly
 100 105 110

Lys Asn Leu His Thr His His Phe Pro Ser Pro Leu Ser Asn Asn Gln
 115 120 125

45 Glu Val Ser Ala Phe Gly Glu Asp Gly Glu Gly Asp Asp Leu Asp Leu
 130 135 140

Trp Thr Val Arg Cys Ser Gly Gln His Trp Glu Arg Glu Ala Ala Val
 145 150 155 160

50 Arg Phe Gln His Val Gly Thr Ser Val Phe Leu Ser Val Thr Gly Glu
 165 170 175

Gln Tyr Gly Ser Pro Ile Arg Gly Gln His Glu Val His Gly Met Pro
 180 185 190

Ser Ala Asn Thr His Asn Thr Trp Lys Ala Met Glu Gly Ile Phe Ile
 195 200 205

60 Lys Pro Ser Val Glu Pro Ser Ala Gly His Asp Glu Leu Xaa

281

210

215

220

5 (2) INFORMATION FOR SEQ ID NO: 136:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 156 amino acids

(B) TYPE: amino acid

10 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 136:

Met Val Ile Glu Ile Ser Asn Lys Thr Ser Ser Ser Ser Thr Cys Ile
1 5 10 15

Leu Val Leu Leu Val Ser Phe Cys Leu Leu Leu Val Pro Ala Met Tyr
20 25 30

Ser Ser Asp Thr Arg Gly Ser Leu Pro Ala Glu His Gly Val Leu Ser
35 40 45

Arg Gln Leu Arg Ala Leu Pro Ser Glu Asp Pro Tyr Gln Leu Glu Leu
50 55 60

Pro Ala Leu Gln Ser Glu Val Pro Lys Asp Ser Thr His Gln Trp Leu
65 70 75 80

Asp Gly Ser Asp Cys Val Leu Gln Ala Pro Gly Asn Thr Ser Cys Leu
85 90 95

Leu His Tyr Met Pro Gln Ala Pro Ser Ala Glu Pro Pro Leu Glu Trp
100 105 110

Pro Phe Pro Asp Leu Phe Ser Glu Pro Leu Cys Arg Gly Pro Ile Leu
115 120 125

Pro Leu Gln Ala Asn Leu Thr Arg Lys Gly Gly Trp Leu Pro Thr Gly
130 135 140

Ser Pro Ser Val Ile Leu Gln Asp Arg Tyr Ser Gly
145 150 155

45 (2) INFORMATION FOR SEQ ID NO: 137:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 233 amino acids

(B) TYPE: amino acid

50 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 137:

Met Met Ile Leu Phe Asn Leu Leu Ile Phe Leu Cys Gly Ala Ala Leu
1 5 10 15

Leu Ala Val Gly Ile Trp Val Ser Ile Asp Gly Ala Ser Phe Leu Lys
20 25 30

Ile Phe Gly Pro Leu Ser Ser Ser Ala Met Gln Phe Val Asn Val Gly
35 40 45

282

Tyr Phe Leu Ile Ala Ala Gly Val Val Val Phe Ala Leu Gly Phe Leu
 50 55 60
 5 Gly Cys Tyr Gly Ala Lys Thr Glu Ser Lys Cys Ala Leu Val Thr Phe
 65 70 75 80
 Phe Phe Ile Leu Leu Leu Ile Phe Ile Ala Glu Val Ala Ala Ala Val
 85 90 95
 10 Val Ala Leu Val Tyr Thr Thr Met Ala Glu His Phe Leu Thr Leu Leu
 100 105 110
 Val Val Pro Ala Ile Lys Lys Asp Tyr Gly Ser Gln Glu Asp Phe Thr
 115 120 125
 15 Gln Val Trp Asn Thr Thr Met Lys Gly Leu Lys Cys Cys Gly Phe Thr
 130 135 140
 20 Asn Tyr Thr Asp Phe Glu Asp Ser Pro Tyr Phe Lys Glu Asn Ser Ala
 145 150 155 160
 Phe Pro Pro Phe Cys Cys Asn Asp Asn Val Thr Asn Thr Ala Asn Glu
 165 170 175
 25 Thr Cys Thr Lys Gln Lys Ala His Asp Gln Lys Val Glu Gly Cys Phe
 180 185 190
 30 Asn Gln Leu Leu Tyr Asp Ile Arg Thr Asn Ala Val Thr Val Gly Gly
 195 200 205
 Val Ala Ala Gly Ile Gly Gly Leu Glu Leu Ala Ala Met Ile Val Ser
 210 215 220
 35 Met Tyr Leu Tyr Cys Asn Leu Gln Xaa
 225 230

40 (2) INFORMATION FOR SEQ ID NO: 138:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 61 amino acids

(B) TYPE: amino acid

45 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 138:

Met Gly Ser Ser Arg Trp Ser Val Ala Cys Pro Thr Gly Leu Gly Val
 1 5 10 15
 50 Leu Met Leu Gly Leu Gly Gly Asp His Pro Pro Gly Ser Gln Val Asp
 20 25 30
 Pro Leu Leu Met Gly Xaa Cys Val Arg Pro Xaa Leu Pro Glu Leu Thr
 35 40 45
 55 Ala Xaa Trp Arg Glu Xaa Gln Xaa Arg Ser Ala Ser Ala
 50 55 60

60

(2) INFORMATION FOR SEQ ID NO: 139:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 73 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 139:

10 Met Gly Trp Leu Phe Leu Lys Val Leu Leu Ala Gly Val Ser Phe Ser
1 5 10 15
Gly Phe Leu Tyr Pro Leu Val Asp Phe Cys Ile Ser Gly Lys Thr Arg
20 25 30
15 Gly Gln Lys Pro Asn Phe Val Ile Ile Leu Ala Asp Asp Met Gly Trp
35 40 45
Gly Asp Trp Gly Ala Asn Trp Ala Glu Thr Lys Asp Thr Ala Asn Leu
20 50 55 60
Asp Lys Met Ala Ser Glu Gly Met Xaa
65 70

25

(2) INFORMATION FOR SEQ ID NO: 140:

(i) SEQUENCE CHARACTERISTICS:

- 30 (A) LENGTH: 377 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 140:

35 Met His Gly Asn Glu Ala Leu Gly Arg Glu Leu Leu Leu Leu Leu Met
1 5 10 15
Gln Phe Leu Cys His Glu Phe Leu Arg Gly Asn Pro Arg Val Thr Arg
20 25 30
40 Leu Leu Ser Glu Met Arg Ile His Leu Leu Pro Ser Met Asn Pro Asp
35 40 45
Gly Tyr Glu Ile Ala Tyr His Arg Gly Ser Glu Leu Val Gly Trp Ala
45 50 55 60
Glu Gly Arg Trp Asn Asn Gln Ser Ile Asp Leu Asn His Asn Phe Ala
65 70 75 80
50 Asp Leu Asn Thr Pro Leu Trp Glu Ala Gln Asp Asp Gly Lys Val Pro
85 90 95
His Ile Val Pro Asn His His Leu Pro Leu Pro Thr Tyr Tyr Thr Leu
100 105 110
55 Pro Asn Ala Thr Val Ala Pro Glu Thr Ala Ala Val Ile Lys Trp Met
115 120 125
Lys Arg Ile Pro Phe Val Leu Ser Ala Asn Leu His Gly Gly Glu Leu
60 130 135 140

284

Val Val Tyr Pro Phe Asp Met Thr Arg Thr Pro Trp Ala Ala Arg
 145 150 155 160
 5 Glu Leu Thr Pro Thr Pro Asp Asp Ala Val Phe Arg Trp Leu Ser Thr
 165 170 175
 Val Tyr Ala Gly Ser Asn Leu Ala Met Gln Asp Thr Ser Arg Arg Pro
 180 185 190
 10 Cys His Ser Gln Asp Phe Ser Val His Gly Asn Ile Ile Asn Gly Ala
 195 200 205
 Asp Trp His Thr Val Pro Gly Ser Met Asn Asp Phe Ser Tyr Leu His
 210 215 220
 15 Thr Asn Cys Phe Glu Val Thr Val Glu Leu Ser Cys Asp Lys Phe Pro
 225 230 235 240
 20 His Glu Asn Glu Leu Pro Gln Glu Trp Glu Asn Asn Lys Asp Ala Leu
 245 250 255
 Leu Thr Tyr Leu Glu Gln Val Arg Met Gly Ile Ala Gly Val Val Arg
 260 265 270
 25 Asp Lys Asp Thr Glu Leu Gly Ile Ala Asp Ala Val Ile Ala Val Asp
 275 280 285
 Gly Ile Asn His Asp Val Thr Thr Ala Trp Gly Gly Asp Tyr Trp Arg
 290 295 300
 30 Leu Leu Thr Pro Gly Asp Tyr Met Val Thr Ala Ser Ala Glu Gly Tyr
 305 310 315 320
 35 His Ser Val Thr Arg Asn Cys Arg Val Thr Phe Glu Glu Gly Pro Phe
 325 330 335
 Pro Cys Asn Phe Val Leu Thr Lys Thr Pro Lys Gln Arg Leu Arg Glu
 340 345 350
 40 Leu Leu Ala Ala Gly Ala Lys Val Pro Pro Asp Leu Arg Arg Arg Leu
 355 360 365
 Glu Arg Leu Arg Gly Gln Lys Asp Xaa
 370 375
 45

(2) INFORMATION FOR SEQ ID NO: 141:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 43 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 141:

Met Ile Cys Leu Ile Leu Leu Leu Gln Ala Val Val Phe Leu Arg Ser
 1 5 10 15

60

Leu His Val Val His Asn Phe Gln Ile Leu Asp Leu Ser Gly Thr Ser

285

20 25 30

Tyr Pro Lys Phe Tyr Gln Thr Leu His Arg Gln
35 40

5

(2) INFORMATION FOR SEQ ID NO: 142:

10 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 41 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 142:

15 Met Val His Val Leu Glu Ile Leu Leu Phe Ile Thr Met Gln Ala Val
1 5 10 15

20 Ser Phe Pro Phe Gln Thr Gln Ile Asp Thr Cys Asn Thr Gln Asp Pro
20 25 30

Ala Glu Arg Gln Pro Ala Ser Ile Val
35 40

25

(2) INFORMATION FOR SEQ ID NO: 143:

30 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 70 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 143:

35 Met Gly Ser Cys Ser Lys Asn Arg Ser Phe Phe Trp Met Thr Gly Leu
1 5 10 15

Leu Val Phe Ile Ser Leu Leu Leu Ser Glu Trp Gln Gly Pro Trp Glu
20 25 30

40 Gly Arg Ala Ile Gly Glu Gly Trp Ala Ser Trp Ala Leu Thr Asn Gly
35 40 45

45 Trp Ala Val Gln Leu Leu Met Ser Leu Gly Asn Asn Thr Glu Lys His
50 55 60

Ser Val Met Ile Tyr Glu
65 70

50

(2) INFORMATION FOR SEQ ID NO: 144:

55 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 483 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 144:

60 Met Ala Thr Gly Gly Gly Ile Arg Ala Met Thr Ser Leu Tyr Gly Gln

286

	1	5	10	15
	Leu Ala Gly	Leu Lys Glu	Leu Gly Leu Leu Asp Cys Xaa	Ser Tyr Ile
		20	25	30
5	Thr Gly Ala	Ser Gly Ser Thr	Trp Ala Leu Ala Asn	Leu Tyr Lys Asp
	35		40	45
	Pro Glu Trp	Ser Gln Lys Asp	Leu Ala Gly Pro Thr	Glu Leu Leu Lys
10	50		55	60
	Thr Gln Val	Thr Lys Asn Lys	Leu Gly Val Leu Ala	Pro Ser Gln Leu
	65	70	75	80
15	Gln Arg Tyr	Arg Gln Glu Leu	Ala Glu Arg Ala Arg	Leu Gly Tyr Pro
		85	90	95
	Ser Cys Phe	Thr Asn Leu Trp	Ala Leu Ile Asn	Glu Ala Leu Leu His
		100	105	110
20	Asp Glu Pro	His Asp His Lys	Leu Ser Asp Gln	Arg Glu Ala Leu Ser
	115		120	125
	His Gly Gln	Asn Pro Leu Pro	Ile Tyr Cys Ala	Leu Asn Thr Lys Gly
25	130		135	140
	Gln Ser Leu	Thr Thr Phe Glu	Phe Gly Glu Trp	Cys Glu Phe Ser Pro
	145	150	155	160
30	Tyr Glu Val	Gly Phe Pro Lys	Tyr Gly Ala Phe	Ile Pro Ser Glu Leu
		165	170	175
	Phe Gly Ser	Glu Phe Phe Met	Gly Gln Leu Met	Lys Arg Leu Pro Glu
		180	185	190
35	Ser Arg Ile	Cys Phe Leu Glu	Gly Ile Trp Ser	Asn Leu Tyr Ala Ala
	195		200	205
	Asn Leu Gln	Asp Ser Leu Tyr	Trp Ala Ser Glu	Pro Ser Gln Phe Trp
40	210		215	220
	Asp Arg Trp	Val Arg Asn Gln	Ala Asn Leu Asp	Lys Glu Gln Val Pro
	225	230	235	240
45	Leu Leu Lys	Ile Glu Glu Pro	Pro Ser Thr Ala	Gly Arg Ile Ala Glu
		245	250	255
	Phe Phe Thr	Asp Leu Leu Thr	Trp Arg Pro Leu	Ala Gln Ala Thr His
		260	265	270
50	Asn Phe Leu	Arg Gly Leu His	Phe His Lys Asp	Tyr Phe Gln His Pro
	275		280	285
	His Phe Ser	Thr Trp Lys Ala	Thr Thr Leu Asn	Gly Leu Pro Asn Gln
55	290		295	300
	Leu Thr Pro	Ser Glu Pro His	Leu Cys Leu Leu	Asp Val Gly Tyr Leu
	305	310	315	320
60	Ile Asn Thr	Ser Cys Leu Pro	Leu Leu Gln Ile	Thr Arg Asp Val Asp

287

[illegible]

35 (2) INFORMATION FOR SEQ ID NO: 145:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 226 amino acids

(B) TYPE: amino acid

40 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 145:

[illegible]

288

Asn Leu Glu Asn Leu Glu Glu Lys Glu Tyr Phe Gly Ile Val Ser Val
 100 105 110
 5 Arg Ile Leu Val His Glu Trp Pro Met Thr Ser Gly Ser Ser Leu Gln
 115 120 125
 Leu Ile Val Ile Gln Glu Glu Val Val Glu Ile Asp Gly Lys Gln Val
 130 135 140
 10 Gln Gln Lys Asp Val Thr Glu Ile Asp Ile Leu Val Lys Asn Arg Gly
 145 150 155 160
 Val Leu Arg His Ser Asn Tyr Thr Leu Pro Leu Glu Glu Ser Met Leu
 165 170 175
 15 Tyr Ser Ile Ser Arg Asp Ser Asp Ile Leu Phe Thr Leu Pro Asn Leu
 180 185 190
 Ser Lys Lys Glu Ser Val Ser Ser Leu Gln Thr Thr Ser Gln Tyr Leu
 20 195 200 205
 Ile Arg Asn Val Glu Thr Thr Val Asp Glu Asp Val Leu Pro Gly Gln
 210 215 220
 25 Val Thr
 225

30 (2) INFORMATION FOR SEQ ID NO: 146:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 45 amino acids
 (B) TYPE: amino acid
 35 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 146:

Met Gly Met Gly Ala Phe Gln Ala Phe Phe Trp Val Ile Leu Thr Val
 40 1 5 10 15
 Ser Asn Val Cys Val Leu Phe Lys Met Ser Leu Phe Phe Leu Leu Thr
 20 25 30
 45 Leu Ile Ser Lys Leu His Gly Asp Ala Glu Val Cys Xaa
 35 40 45

50 (2) INFORMATION FOR SEQ ID NO: 147:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 132 amino acids
 (B) TYPE: amino acid
 55 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 147:

Met Ser Gly Gly Trp Met Ala Gln Val Gly Ala Trp Arg Thr Gly Ala
 1 5 10 15
 60 Leu Gly Leu Ala Leu Leu Leu Leu Leu Gly Leu Gly Leu Gly Leu Glu

289

20 25 30

Ala Pro Arg Ala Arg Phe Pro Pro Arg Pro Leu Pro Arg Pro His Pro
35 40 45

5 Ser Ser Gly Ser Cys Pro Pro Thr Lys Phe Gln Cys Arg Thr Ser Gly
50 55 60

10 Leu Cys Val Pro Leu Thr Trp Arg Cys Asp Arg Thr Trp Thr Ala Ala
65 70 75 80

Met Ala Ala Met Arg Arg Ser Ala Gly Leu Ser His Val Pro Arg Lys
85 90 95

15 Gly Asn Ala His Arg Pro Leu Ala Ser Pro Ala Pro Ala Pro Ala Ser
100 105 110

Val Thr Ala Leu Gly Glu Leu Thr Arg Asn Cys Ala Thr Ala Ala Ala
115 120 125

20 Trp Pro Ala Xaa
130

25

(2) INFORMATION FOR SEQ ID NO: 148:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 92 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 148:

30

35 Met Glu Ala Thr Leu Glu Gln His Leu Glu Asp Thr Met Lys Asn Pro
1 5 10 15

Ser Ile Val Gly Val Leu Cys Thr Asp Ser Gln Gly Leu Asn Leu Gly
20 25 30

40 Cys Arg Gly Thr Leu Ser Asp Glu His Ala Gly Val Ile Ser Val Leu
35 40 45

Ala Gln Gln Ala Ala Lys Leu Thr Ser Asp Pro Thr Asp Ile Pro Val
50 55 60

45 Val Cys Leu Glu Ser Asp Asn Gly Asn Ile Met Ile Gln Lys His Asp
65 70 75 80

50 Gly Ile Thr Val Ala Val His Lys Met Ala Ser Xaa
85 90

55

(2) INFORMATION FOR SEQ ID NO: 149:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 165 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 149:

60

290

Met Glu Pro Leu Arg Leu Leu Ile Leu Leu Phe Val Thr Glu Leu Ser
 1 5 10 15

5 Gly Ala His Asn Thr Thr Val Phe Gln Gly Val Ala Gly Gln Ser Leu
 20 25 30

Gln Val Ser Cys Pro Tyr Asp Ser Met Lys His Trp Gly Arg Arg Lys
 35 40 45

10 Ala Trp Cys Arg Gln Leu Gly Glu Lys Gly Pro Cys Gln Arg Val Val
 50 55 60

15 Ser Thr His Asn Leu Trp Leu Leu Ser Phe Leu Arg Arg Trp Asn Gly
 65 70 75 80

Ser Thr Ala Ile Thr Asp Asp Thr Leu Gly Gly Thr Leu Thr Ile Thr
 85 90 95

20 Leu Arg Asn Leu Gln Pro His Asp Ala Gly Leu Tyr Gln Cys Gln Ser
 100 105 110

Leu His Gly Ser Glu Ala Asp Thr Leu Arg Lys Val Leu Val Glu Val
 115 120 125

25 Leu Ala Asp Pro Leu Asp His Arg Asp Ala Gly Asp Leu Trp Phe Pro
 130 135 140

30 Gly Glu Ser Glu Ser Phe Glu Asp Ala His Val Glu His Ser Ile Ser
 145 150 155 160

Arg Ser Ser Ser Xaa
 165

35

(2) INFORMATION FOR SEQ ID NO: 150:

40

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 139 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 150:

45

Met Ile Ser Leu Thr Asp Thr Gln Lys Ile Gly Met Gly Leu Thr Gly
 1 5 10 15

Phe Gly Val Phe Phe Leu Phe Phe Gly Met Ile Leu Phe Phe Asp Lys
 20 25 30

50

Ala Leu Leu Ala Ile Gly Asn Val Leu Phe Val Ala Gly Leu Ala Phe
 35 40 45

55

Val Ile Gly Leu Glu Arg Thr Phe Arg Phe Phe Phe Gln Lys His Lys
 50 55 60

Met Lys Ala Thr Gly Phe Phe Leu Gly Gly Val Phe Val Val Leu Ile
 65 70 75 80

60

Gly Trp Pro Leu Ile Gly Met Ile Phe Glu Ile Tyr Gly Phe Phe Leu

.291

85 90 95

Leu Phe Arg Gly Phe Phe Pro Val Val Val Gly Phe Ile Arg Arg Val
100 105 110

5 Pro Val Leu Gly Ser Leu Leu Asn Leu Pro Gly Ile Arg Ser Phe Val
115 120 125

10 Asp Lys Val Gly Glu Ser Asn Asn Met Val Xaa
130 135

15 (2) INFORMATION FOR SEQ ID NO: 151:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 58 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 151:

Met Ser Ala Pro Gln Thr Arg Ile Ser Arg Ala Leu Val Leu Leu Phe
1 5 10 15

25 Leu Ala Pro Thr Leu Leu Ser Leu Gly His Gly Ile His Pro Ile Asn
20 25 30

Thr Ala Thr Pro Tyr Xaa Thr Asp Gln Ala Lys Leu Ala Pro Gly Thr
35 40 45

30 Lys Glu Leu Asn His Asp Gln Ser Val Thr
50 55

35

(2) INFORMATION FOR SEQ ID NO: 152:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 48 amino acids

40 (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 152:

Met Ile Arg Lys Leu His Lys Ile Ile Val Phe Ser Pro Arg Val Ile
1 5 10 15

Val Leu Leu Asn Cys Phe Phe Phe Ile Lys Ala Lys Phe Val Leu Tyr
20 25 30

50 Ile Phe Val Phe His Val Leu Asp Gly Ser Ile Ser Tyr Pro Val Xaa
35 40 45

55

(2) INFORMATION FOR SEQ ID NO: 153:

60

(i) SEQUENCE CHARACTERISTICS:

292

(A) LENGTH: 42 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 153:

5

Met Leu Leu Asn Gln His Phe Lys Ile Phe Gly Ser Leu Ile His Met
 1 5 10 15

10

Asn Leu Leu Phe Ala Leu Ile Ser Leu Gly Ser Ser Asn Leu Ser Gly
 20 25 30

Val Gln Phe Cys Cys Glu Thr Val Gln Xaa
 35 40

15

(2) INFORMATION FOR SEQ ID NO: 154:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 72 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 154:

20

25

Met Leu Ser Leu Ser Phe Leu Leu Arg Arg Val Leu Phe Leu Gly Phe
 1 5 10 15

Leu Gln Ala Ser Val Gly Glu Lys Lys Ser Leu Arg Xaa Leu Asn Tyr
 20 25 30

30

Ser Val Pro His Pro Met Leu Xaa His Pro Pro Pro Asp Thr Ala Gln
 35 40 45

35

Val Pro Pro Arg Leu Glu Arg Ser Leu Leu Gln Gln Glu Leu Trp Thr
 50 55 60

Pro Gly Pro His His Ser Asn Ile
 65 70

40

(2) INFORMATION FOR SEQ ID NO: 155:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 106 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 155:

45

50

Met Gln Pro Leu Asn Phe Ser Ser Thr Glu Cys Ser Ser Phe Ser Pro
 1 5 10 15

Pro Thr Thr Val Ile Leu Leu Ile Leu Leu Cys Phe Glu Gly Leu Leu
 20 25 30

55

Phe Leu Ile Phe Thr Ser Val Met Phe Gly Thr Gln Val His Ser Ile
 35 40 45

60

Cys Thr Asp Glu Thr Gly Ile Glu Gln Leu Lys Lys Glu Glu Arg Arg
 50 55 60

293

Trp Ala Lys Lys Thr Lys Trp Met Asn Met Lys Ala Val Phe Gly His
65 70 75 80

5 Pro Phe Ser Leu Gly Trp Ala Ser Pro Phe Ala Thr Pro Asp Gln Gly
85 90 95

Lys Ala Asp Pro Tyr Gln Tyr Val Val Xaa
100 105

10

(2) INFORMATION FOR SEQ ID NO: 156:

15

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 29 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 156:

20

Met Tyr Thr Asn His Phe Asn Leu Tyr Leu Lys Tyr Ile Leu Leu Ile
1 5 10 15

25

Ile Leu Ile Leu Asn Met Thr Asn Ser Ser Ser Arg Tyr
20 25

(2) INFORMATION FOR SEQ ID NO: 157:

30

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 53 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 157:

Met Asn Glu Leu Leu Leu Phe Phe Phe Phe Phe Phe Phe Thr Phe
1 5 10 15

40

Cys Ile Glu Thr Asn Ser Phe Lys Gln Thr Tyr Tyr Tyr Tyr Phe Leu
20 25 30

Gln Asn Ile Tyr Met Glu Met Leu Pro Pro Pro Val Asn Pro Pro Val
35 40 45

45

Pro Pro Trp Gly Xaa
50

50

(2) INFORMATION FOR SEQ ID NO: 158:

55

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 75 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 158:

60

Met Tyr Ala Val Tyr Gln Gln Leu Ala Gln Leu Thr Leu Met Val Thr
1 5 10 15

	Leu	Leu	Ala	Pro	Ile	Leu	Pro	Asp	Glu	Gln	Ser	Glu	Val	Phe	Glu	Ala
				20					25					30		
5	Leu	Ser	Asn	Leu	Pro	Lys	Val	Thr	Trp	Leu	Gly	Ser	Asn	Ser	Pro	Ser
			35					40					45			
	Ser	Glu	Met	Pro	Glu	Pro	Gly	Arg	Phe	Val	Ile	Val	His	His	Gln	Leu
			50				55					60				
10	Ser	Ala	Ala	Ser	His	Ser	Ser	Ser	Gln	Leu	Ala					
		65				70					75					

15

(2) INFORMATION FOR SEQ ID NO: 159:

20 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 81 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 159:

25	Met Trp Pro Pro Leu Leu Leu Leu Leu Leu Pro Ala Ala Pro 1 5 10 15
	Val Pro Thr Ala Lys Ala Ala Pro His Pro Asp Ala Asn Thr Gln Glu 20 25 30
30	Gly Leu Gln Asn Leu Leu Gln Gly Val Gly Ala Gly Gly Asp Gly Glu 35 40 45
	Leu Arg Ala Asp Ser His Leu Ala Pro Gly Ser Gly Cys Ile Asp Gly 50 55 60
35	Ala Val Val Ala Thr Arg Pro Glu Ser Arg Gly Gly Arg Pro Ala Val 65 70 75 80
40	Pro

45 (2) INFORMATION FOR SEQ ID NO: 160:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 139 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 160:

Met Lys Phe Thr Thr Leu Leu Phe Leu Ala Ala Val Ala Gly Ala Leu
1 5 10 15

55 Val Tyr Ala Glu Asp Ala Ser Ser Asp Ser Thr Gly Ala Asp Pro Ala
20 25 30

Gln Glu Ala Gly Thr Ser Lys Pro Asn Glu Glu Ile Ser Gly Pro Ala
35 40 45

60

295

Glu Pro Ala Ser Pro Pro Glu Thr Thr Thr Thr Ala Gln Glu Thr Ser
 50 55 60

5 Ala Ala Ala Val Gln Gly Thr Ala Lys Val Thr Ser Ser Arg Gln Glu
 65 70 75 80

Leu Asn Pro Leu Lys Ser Ile Val Glu Lys Ser Ile Leu Leu Thr Glu
 85 90 95

10 Gln Ala Leu Ala Lys Ala Gly Lys Gly Met His Gly Gly Val Pro Gly
 100 105 110

Gly Lys Gln Phe Ile Glu Asn Gly Ser Glu Phe Ala Gln Lys Leu Leu
 115 120 125

15 Lys Lys Phe Ser Leu Leu Lys Pro Trp Ala Xaa
 130 135

20 (2) INFORMATION FOR SEQ ID NO: 161:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 178 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 161:

30 Met Leu Gly Cys Gly Ile Pro Ala Leu Gly Leu Leu Leu Leu Leu Gln
 1 5 10 15

Gly Ser Ala Asp Gly Asn Gly Ile Gln Gly Phe Phe Tyr Pro Trp Ser
 20 25 30

35 Cys Glu Gly Asp Ile Trp Asp Arg Glu Ser Cys Gly Gly Gln Ala Ala
 35 40 45

Ile Asp Ser Pro Asn Leu Cys Leu Arg Leu Arg Cys Cys Tyr Arg Asn
 50 55 60

40 Gly Val Cys Tyr His Gln Arg Pro Asp Glu Asn Val Arg Arg Lys His
 65 70 75 80

45 Met Trp Ala Leu Val Trp Thr Cys Ser Gly Leu Leu Leu Leu Ser Cys
 85 90 95

Ser Ile Cys Leu Phe Trp Trp Ala Lys Arg Arg Asp Val Leu His Met
 100 105 110

50 Pro Gly Phe Leu Ala Gly Pro Cys Asp Met Ser Lys Ser Val Ser Leu
 115 120 125

Leu Ser Lys His Arg Gly Thr Lys Lys Thr Thr Thr Thr Gly Ser Val
 130 135 140

55 Pro Val Ala Leu Ser Lys Glu Ser Arg Arg Arg Arg Gly Gly Thr Glu
 145 150 155 160

60 Gly Glu Gly Thr Glu Glu Gly Glu Glu Thr Thr Thr Thr Thr Thr Thr
 165 170 175

Asp Xaa

5

(2) INFORMATION FOR SEQ ID NO: 162:

10

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 72 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 162:

15

Met Glu Ala Val Phe Thr Val Phe Phe Phe Val Val Val Leu Phe Leu
1 5 10 15

Lys Asn Thr Glu Gly Ala Lys Leu Phe Cys Thr Leu Tyr Pro Ala Ala
20 25 30

20

Ser Ser Gly Gln Ser Gln Gly Pro Gly Leu Glu Lys Pro Asp Ser Gln
35 40 45

25

Glu Cys Ile Ile Asp Pro Cys Ser Tyr Pro Ile Ala Leu Gly Ala Gly
50 55 60

Thr Glu Pro Gly Cys Lys Ile Xaa
65 70

30

(2) INFORMATION FOR SEQ ID NO: 163:

35

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 67 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 163:

40

Met Trp Phe Tyr Phe Leu Ser Val Ser Phe Pro Leu Leu Pro Val Xaa
1 5 10 15

Ala Pro Xaa Pro Pro Pro Ala Pro Thr Thr Leu Cys Leu Leu Leu Phe
20 25 30

45

Leu Gly Xaa Leu Tyr Asn Ser Thr Cys Ile His Cys Val His Thr Thr
35 40 45

50

Ser Xaa Thr Gln Asn Pro Thr Ala Asn Thr Leu Lys Lys Lys Lys Lys
50 55 60

Asn Trp Gly
65

55

(2) INFORMATION FOR SEQ ID NO: 164:

60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 155 amino acids

297

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 164:

5 Met Gly Phe Gly Ala Thr Leu Ala Val Gly Leu Thr Ile Phe Val Leu
 1 5 10 15

Ser Val Val Thr Ile Ile Ile Cys Phe Thr Cys Ser Cys Cys Cys Leu
 20 25 30

10 Tyr Lys Thr Cys Arg Arg Pro Arg Pro Val Val Thr Thr Thr Thr Ser
 35 40 45

Thr Thr Val Val His Ala Pro Tyr Pro Gln Pro Pro Ser Val Pro Pro
 15 50 55 60

Ser Tyr Pro Gly Pro Ser Tyr Gln Gly Tyr His Thr Met Pro Pro Gln
 65 70 75 80

20 Pro Gly Met Pro Ala Ala Pro Tyr Pro Met Gln Tyr Pro Pro Pro Tyr
 85 90 95

Pro Ala Gln Pro Met Gly Pro Pro Ala Tyr His Glu Thr Leu Ala Gly
 100 105 110

25 Glu Gln Pro Arg Pro Thr Pro Pro Ala Ser Leu Leu Thr Thr Arg Pro
 115 120 125

Thr Trp Met Pro Arg Arg Arg Pro Ser Glu His Ser Leu Ala Ser Leu
 130 135 140

Ala Ala Thr Trp Leu Cys Cys Val Cys Ala Xaa
 145 150 155

35

(2) INFORMATION FOR SEQ ID NO: 165:

(i) SEQUENCE CHARACTERISTICS:

40 (A) LENGTH: 104 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 165:

45 Met Ile Ile Leu Val Phe Ile Ala Phe Phe Ile Pro Leu Gln Lys Thr
 1 5 10 15

Ile Gly Lys Ile Ala Thr Cys Leu Glu Leu Arg Ser Ala Ala Leu Gln
 20 25 30

50 Ser Thr Gln Ser Gln Glu Glu Phe Lys Leu Glu Asp Leu Lys Lys Leu
 35 40 45

Glu Pro Ile Leu Lys Asn Ile Leu Thr Tyr Asn Lys Glu Phe Pro Phe
 55 60

Asp Val Gln Pro Val Pro Leu Arg Arg Ile Leu Ala Pro Gly Glu Glu
 65 70 75 80

60 Glu Asn Leu Glu Phe Glu Glu Asp Glu Glu Glu Gly Gly Ala Gly Ala

298

85

90

95

Gly Leu Leu Ile Leu Ser Cys Xaa
100

5

(2) INFORMATION FOR SEQ ID NO: 166:

10

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 81 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 166:

15

Met Ala Gly Thr Met Val Ile Val Val Val Val Val Gly Glu Val
1 5 10 15

20

Val Val Glu Ala Glu Val Val Val Gln Ala Arg Glu Glu Ala Gly Glu
20 25 30

Glu Glu Gly Ala Arg Ile Ile Thr Lys Gly Val Asn Leu Asn Ser Ile
35 40 45

25

Ser Ser Met Glu Val Ile Ser Ile Ile Ile Leu Asp Leu Asp Arg Glu
50 55 60

Asp Ile Thr Leu Val Glu Ala Thr Glu Pro Tyr Ile Leu Leu Glu Leu
65 70 75 80

30

Lys

35

(2) INFORMATION FOR SEQ ID NO: 167:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 93 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 167:

40

Met Ser Phe Ser Phe Ile Ile Phe Leu Leu Leu Val Cys Gln Glu Ile
1 5 10 15

45

Thr Phe Cys Met Ser Tyr Gly Asp Ala Val Asn Cys Phe Ser Glu Cys
20 25 30

50

Phe Ser Asn Leu Gln Thr Ile Tyr Ile Ser Cys Leu Gln His Ala Val
35 40 45

Cys Lys His Ser Val Ile Trp Ser Ile Gln Leu Phe Val Arg Ala Leu
50 55 60

55

Pro Ile Ser Lys Cys Ala Glu Leu Ser Ile Asp Gly Ile Phe Arg Ser
65 70 75 80

60

Phe His Glu Asn Trp Lys Cys Ser Trp Val Ala Pro Thr
85 90

5 (2) INFORMATION FOR SEQ ID NO: 168:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 58 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 168:

Met Gly Trp Ser Ala Gly Leu Leu Phe Leu Leu Ile Leu Tyr Leu Pro
1 5 10 15

15 Val Pro Gly Trp Met Glu Arg Glu Asp Gly Glu Thr Gly His Leu Ser
20 25 30

Pro Gln Ala Pro Gly Arg Glu Tyr Arg Gly Phe Tyr Ser Val Pro Pro
35 40 45

20 Asp Tyr Val Trp Leu Arg Asp Ser Pro Xaa
50 55

25 (2) INFORMATION FOR SEQ ID NO: 169:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 232 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 169:

Met Ala Thr Leu Trp Gly Gly Leu Leu Arg Leu Gly Ser Leu Leu Ser
35 1 5 10 15

Leu Ser Cys Leu Ala Leu Ser Val Leu Leu Ala His Cys Gln Thr
20 25 30

40 Pro Pro Arg Ile Ser Arg Met Ser Asp Val Asn Val Ser Ala Leu Pro
35 40 45

Ile Lys Lys Asn Ser Gly His Ile Tyr Asn Lys Asn Ile Ser Gln Lys
50 55 60

45 Asp Cys Asp Cys Leu His Val Val Glu Pro Met Pro Val Arg Gly Pro
65 70 75 80

50 Asp Val Glu Ala Tyr Cys Leu Arg Cys Glu Cys Lys Tyr Glu Glu Arg
85 90 95

Ser Ser Val Thr Ile Lys Val Thr Ile Ile Ile Tyr Leu Ser Ile Leu
100 105 110

55 Gly Leu Leu Leu Leu Tyr Met Val Tyr Leu Thr Leu Val Glu Pro Ile
115 120 125

Leu Lys Arg Arg Leu Phe Gly His Ala Gln Leu Ile Gln Ser Asp Asp
130 135 140

60

300

Asp Ile Gly Asp His Gln Pro Phe Ala Asn Ala His Asp Val Leu Ala
 145 150 155 160
 5 Arg Ser Arg Ser Arg Ala Asn Val Leu Asn Lys Val Glu Tyr Gly Thr
 165 170 175
 Ala Ala Leu Glu Ala Ser Ser Pro Arg Ala Ala Lys Ser Leu Ser Leu
 180 185 190
 10 Thr Gly Met Leu Ser Ser Ala Asn Trp Gly Ile Glu Phe Lys Val Thr
 195 200 205
 Arg Lys Lys Gln Ala Asp Asn Trp Lys Gly Thr Asp Trp Val Leu Leu
 210 215 220
 15 Gly Phe Ile Leu Ile Pro Cys Xaa
 225 230

20

(2) INFORMATION FOR SEQ ID NO: 170:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 72 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 170:

30 Met Ser Ala Ile Phe Asn Phe Gln Ser Leu Leu Thr Val Ile Leu Leu
 1 5 10 15
 Leu Ile Cys Thr Cys Ala Tyr Ile Arg Ser Leu Ala Pro Ser Leu Leu
 20 25 30
 35 Asp Arg Asn Lys Thr Gly Leu Leu Gly Ile Phe Trp Lys Cys Ala Arg
 35 40 45
 Ile Gly Glu Arg Lys Ser Pro Tyr Val Ala Val Cys Cys Ile Val Met
 50 55 60
 40 Ala Phe Ser Ile Leu Phe Ile Gln
 65 70

45

(2) INFORMATION FOR SEQ ID NO: 171:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 65 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 171:

55 Met Gly Thr Phe Ser Leu Ser Leu Phe Gly Leu Met Gly Val Ala Phe
 1 5 10 15
 Gly Met Asn Leu Glu Ser Ser Leu Glu Glu Asp His Arg Ile Phe Trp
 20 25 30
 60 Leu Ile Thr Gly Ile Met Phe Met Gly Ser Gly Leu Ile Trp Arg Arg

35 40 45

Leu Leu Ser Phe Leu Gly Arg Gln Leu Glu Ala Pro Leu Pro Pro Met
50 55 60

Val
65

(2) INFORMATION FOR SEQ ID NO: 172:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 75 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 172:

Met Tyr Lys Gly Lys Leu Val Ile Val Leu Ile Leu Leu Leu Leu Pro
1 5 10 15

Ser His Phe Met Phe Leu Thr Gln Cys Lys Glu Ile Lys His Asn Leu
20 25 30

Lys Lys Asn Met Ser Leu Leu Leu Phe Thr Ile Lys Ser Trp Leu Tyr
25 35 40 45

Ser Ala Ser Leu Gly Ile Leu Tyr Asn Trp Gln His Leu Thr Ala Gln
50 55 60

Val Asp Gln Cys Thr Ser Leu Ile Leu Ile His
65 70 75

(2) INFORMATION FOR SEQ ID NO: 173:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 334 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 173:

Met Val Gly His Glu Met Ala Ser Xaa Ser Ser Asn Thr Ser Leu Pro
1 5 10 15

Phe Ser Asn Met Gly Asn Pro Met Asn Thr Thr Gln Leu Gly Lys Ser
20 25 30

Leu Phe Gln Trp Gln Val Glu Gln Glu Glu Ser Lys Leu Ala Asn Ile
35 40 45

Ser Gln Asp Gln Phe Leu Ser Lys Asp Ala Asp Gly Asp Thr Phe Leu
50 55 60

His Ile Ala Val Ala Gln Gly Arg Arg Ala Leu Ser Tyr Val Leu Ala
65 70 75 80

Arg Lys Met Asn Ala Leu His Met Leu Asp Ile Lys Glu His Asn Gly
85 90 95

302

Gln Ser Ala Phe Gln Val Ala Val Ala Ala Asn Gln His Leu Ile Val
 100 105 110
 5 Gln Asp Leu Val Asn Ile Gly Ala Gln Val Asn Thr Thr Asp Cys Trp
 115 120 125
 Gly Arg Thr Pro Leu His Val Cys Ala Glu Lys Gly His Ser Gln Val
 130 135 140
 10 Leu Gln Ala Ile Gln Lys Gly Ala Val Gly Ser Asn Gln Phe Val Asp
 145 150 155 160
 Leu Glu Ala Thr Asn Tyr Asp Gly Leu Thr Pro Leu His Cys Ala Val
 165 170 175
 15 Ile Ala His Asn Ala Val Val His Glu Leu Gln Arg Asn Gln Gln Pro
 180 185 190
 20 His Ser Pro Glu Val Gln Glu Leu Leu Leu Lys Asn Lys Ser Leu Val
 195 200 205
 Asp Thr Ile Lys Cys Leu Ile Gln Met Gly Ala Ala Val Glu Ala Lys
 210 215 220
 25 Asp Arg Lys Ser Gly Arg Thr Ala Leu His Leu Ala Ala Glu Glu Ala
 225 230 235 240
 Asn Leu Glu Leu Ile Arg Leu Phe Leu Glu Leu Pro Ser Cys Leu Ser
 245 250 255
 Phe Val Asn Ala Lys Ala Tyr Asn Gly Asn Thr Ala Leu His Val Ala
 260 265 270
 35 Ala Ser Leu Gln Tyr Arg Leu Thr Gln Leu Asp Ala Val Arg Leu Leu
 275 280 285
 Met Arg Lys Gly Ala Asp Pro Ser Thr Arg Asn Leu Glu Asn Glu Gln
 290 295 300
 40 Pro Val His Leu Val Pro Asp Gly Pro Val Gly Glu Gln Ile Arg Arg
 305 310 315 320
 Ile Leu Lys Gly Lys Ser Ile Gln Gln Arg Ala Pro Pro Tyr
 325 330
 45

(2) INFORMATION FOR SEQ ID NO: 174:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 196 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 174:

Met Asp Ala Arg Trp Trp Ala Val Val Val Leu Ala Ala Phe Pro Ser
 1 5 10 15

60 Leu Gly Ala Gly Gly Glu Thr Pro Glu Ala Pro Pro Glu Ser Trp Thr

303

[illegible]

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(2) INFORMATION FOR SEQ ID NO: 175:

40      (i) SEQUENCE CHARACTERISTICS:
          (A) LENGTH: 265 amino acids
          (B) TYPE: amino acid
          (D) TOPOLOGY: linear

          (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 175:

45      Met Ser Asp Leu Leu Leu Leu Gly Leu Ile Gly Gly Leu Thr Leu Leu
          1             5             10             15

          Leu Leu Leu Thr Leu Leu Ala Phe Ala Gly Tyr Ser Gly Leu Leu Ala
50          20             25             30

          Gly Val Glu Val Ser Ala Gly Ser Pro Pro Ile Arg Asn Val Thr Val
          35             40             45

          Ala Tyr Lys Phe His Met Gly Leu Tyr Gly Glu Thr Gly Arg Leu Phe
55          50             55             60

          Thr Glu Ser Cys Ser Ile Ser Pro Lys Leu Arg Ser Ile Ala Val Tyr
          65             70             75             80
60

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304

Tyr Asp Asn Pro His Met Val Pro Pro Asp Lys Cys Arg Cys Ala Val
 85 90 95
 5 Gly Ser Ile Leu Ser Glu Gly Glu Glu Ser Pro Ser Pro Glu Leu Ile
 100 105 110
 Asp Leu Tyr Gln Lys Phe Gly Phe Lys Val Phe Ser Phe Pro Glu Pro
 115 120 125
 10 Ser His Val Val Thr Ala Thr Phe Pro Leu Thr Pro Pro Phe Cys Pro
 130 135 140
 Ile Trp Leu Gly Tyr Pro Pro Cys Pro Ser Cys Leu Gly His Leu His
 15 145 150 155 160
 Gln Gly Ala Glu Ala Val Cys Leu Ser Ser Ala Gly Asp Leu Pro Gly
 165 170 175
 20 Arg Pro Glu Ser Ile Ser Cys Ala His Trp His Gly Gln Gly Asp Phe
 180 185 190
 Tyr Val Pro Glu Met Lys Glu Thr Glu Trp Lys Trp Arg Gly Leu Val
 195 200 205
 25 Glu Ala Ile Asp Thr Gln Val Asp Gly Thr Gly Ala Asp Thr Met Ser
 210 215 220
 Asp Thr Ser Ser Val Ser Leu Glu Val Ser Pro Gly Ser Arg Glu Thr
 30 225 230 235 240
 Ser Ala Ala Thr Leu Ser Pro Gly Ala Ser Ser Arg Gly Trp Asp Asp
 245 250 255
 35 Gly Asp Thr Arg Ser Glu His Ser Xaa
 260 265
 (2) INFORMATION FOR SEQ ID NO: 176:
 40 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 138 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 176:
 Met Ala Gln Leu Phe Leu Pro Leu Leu Ala Ala Leu Val Leu Ala Gln
 1 5 10 15
 50 Ala Pro Ala Ala Leu Ala Asp Val Leu Glu Gly Asp Ser Ser Glu Asp
 20 25 30
 Arg Ala Phe Arg Val Arg Ile Ala Gly Asp Ala Pro Leu Gln Gly Val
 35 40 45
 55 Leu Gly Gly Ala Leu Thr Ile Pro Cys His Val His Tyr Leu Arg Pro
 50 55 60
 60 Pro Pro Ser Arg Arg Ala Val Leu Gly Ser Pro Arg Val Lys Trp Thr
 65 70 75 80

305

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Phe Leu Ser Arg Gly Arg Glu Ala Glu Val Leu Val Ala Arg Gly Val
      85                                90                                95
5  Arg Val Lys Val Asn Glu Ala Tyr Arg Phe Arg Val Ala Leu Pro Ala
      100                                105                                110
Tyr Pro Ala Ser Leu Thr Asp Val Ser Pro Gly Ala Glu Arg Ala Ala
      115                                120                                125
10 Pro Gln Arg Leu Arg Tyr Leu Ser Leu Xaa
      130                                135
15
(2) INFORMATION FOR SEQ ID NO: 177:
      (i) SEQUENCE CHARACTERISTICS:
            (A) LENGTH: 179 amino acids
            (B) TYPE: amino acid
            (D) TOPOLOGY: linear
      (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 177:
25 Met Pro Ala Leu Arg Pro Ala Leu Leu Trp Ala Leu Leu Ala Leu Trp
      1          5          10          15
Leu Cys Cys Ala Thr Pro Ala His Ala Leu Gln Cys Arg Asp Gly Tyr
      20          25          30
30 Glu Pro Cys Val Asn Glu Gly Met Cys Val Thr Tyr His Asn Gly Thr
      35          40          45
Gly Tyr Cys Lys Gly Pro Glu Gly Phe Leu Gly Glu Tyr Cys Gln His
      50          55          60
35 Arg Asp Pro Cys Glu Lys Asn Arg Cys Gln Asn Gly Gly Thr Cys Val
      65          70          75          80
Ala Gln Ala Met Leu Gly Lys Ala Thr Cys Arg Cys Ala Ser Gly Phe
      85          90          95
40 Thr Gly Glu Asp Cys Gln Tyr Ser Thr Ser His Pro Cys Phe Val Ser
      100          105          110
45 Arg Pro Cys Leu Asn Gly Gly Thr Cys His Met Leu Ser Arg Asp Thr
      115          120          125
Tyr Glu Cys Thr Cys Gln Val Gly Phe Thr Gly Lys Glu Cys Gln Trp
      130          135          140
50 Thr Asp Ala Cys Leu Ser His Pro Cys Ala Asn Gly Ser Thr Cys Thr
      145          150          155          160
55 Thr Val Ala Asn His Phe Leu Gln Met Pro Phe Ala Leu His Arg Ala
      165          170          175
Glu Val Xaa

```

60

(2) INFORMATION FOR SEQ ID NO: 178:

(i) SEQUENCE CHARACTERISTICS:

5

(A) LENGTH: 155 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 178:

10 Met Thr Arg Gly Gly Pro Gly Gly Arg Pro Gly Leu Pro Gln Pro Pro
1 5 10 15

Pro Leu Leu Leu Leu Leu Leu Pro Leu Leu Leu Val Thr Ala Glu
20 25 30

15 Pro Pro Lys Pro Ala Gly Val Tyr Tyr Ala Thr Ala Tyr Trp Met Pro
35 40 45

Ala Glu Lys Thr Val Gln Val Lys Asn Val Met Asp Lys Asn Gly Asp
20 50 55 60

Ala Tyr Gly Phe Tyr Asn Asn Ser Val Lys Thr Thr Gly Trp Gly Ile
65 70 75 80

25 Leu Glu Ile Arg Ala Gly Tyr Gly Ser Gln Thr Leu Ser Asn Glu Ile
85 90 95

Ile Met Phe Val Ala Gly Phe Leu Glu Gly Tyr Leu Ile Ala Pro His
100 105 110

30 Met Asn Asp His Tyr Thr Asn Leu Tyr Pro Gln Leu Ile Thr Lys Pro
115 120 125

Ser Ile Met Asp Lys Val Gln Asp Phe Met Glu Lys Gln Asp Lys Val
35 130 135 140

Asp Pro Glu Lys Tyr Gln Arg Ile Gln Asp Xaa
145 150 155

40

(2) INFORMATION FOR SEQ ID NO: 179:

(i) SEQUENCE CHARACTERISTICS:

45

(A) LENGTH: 295 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 179:

50 Met Leu Gln Gly Pro Gly Ser Leu Leu Leu Leu Phe Leu Ala Ser His
1 5 10 15

Cys Cys Leu Gly Ser Ala Arg Gly Leu Phe Leu Phe Gly Gln Pro Asp
20 25 30

55 Phe Ser Tyr Lys Arg Xaa Asn Cys Lys Pro Leu Phe Val Asn Leu Gln
35 40 45

Leu Cys His Gly Ile Glu Tyr Gln Asn Met Asn Leu Pro Asn Leu Leu
60 50 55 60

307

Gly His Glu Thr Met Lys Glu Val Leu Glu Gln Ala Gly Ala Trp Ile
 65 70 75 80
 5 Pro Leu Val Met Lys Gln Cys His Pro Asp Thr Lys Lys Phe Leu Cys
 85 90 95
 Ser Leu Phe Ala Pro Val Cys Leu Asp Asp Leu Asp Glu Thr Ile Gln
 100 105 110
 10 Pro Cys His Ser Leu Cys Val Gln Val Lys Asp Arg Cys Ala Pro Val
 115 120 125
 Met Ser Ala Phe Gly Phe Pro Trp Pro Asp Met Leu Glu Cys Asp Arg
 130 135 140
 15 Phe Pro Gln Asp Asn Asp Leu Cys Ile Pro Leu Ala Ser Ser Asp His
 145 150 155 160
 20 Leu Leu Pro Ala Thr Glu Glu Ala Pro Lys Val Cys Glu Ala Cys Lys
 165 170 175
 Asn Lys Asn Asp Asp Asp Asn Asp Ile Met Glu Thr Leu Cys Lys Asn
 180 185 190
 25 Asp Phe Ala Leu Lys Ile Lys Val Lys Glu Ile Thr Tyr Ile Asn Arg
 195 200 205
 Asp Thr Lys Ile Ile Leu Glu Thr Lys Ser Lys Thr Ile Tyr Lys Leu
 210 215 220
 30 Asn Gly Val Ser Glu Arg Asp Leu Lys Lys Ser Val Leu Trp Leu Lys
 225 230 235 240
 35 Asp Ser Leu Gln Cys Thr Cys Glu Glu Met Asn Asp Ile Asn Ala Pro
 245 250 255
 Tyr Leu Val Met Gly Gln Lys Gln Gly Gly Glu Leu Val Ile Thr Ser
 260 265 270
 40 Val Lys Arg Trp Gln Lys Gly Gln Arg Glu Phe Lys Arg Ile Ser Arg
 275 280 285
 Ser Ile Arg Lys Leu Gln Cys
 290 295
 45

(2) INFORMATION FOR SEQ ID NO: 180:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 256 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 180:

Met Arg Pro Ala Ala Leu Arg Gly Ala Leu Leu Gly Cys Leu Cys Leu
 1 5 10 15
 60 Ala Leu Leu Cys Leu Gly Gly Ala Asp Lys Arg Leu Arg Asp Asn His

308

20 25 30
 Glu Trp Lys Lys Leu Ile Met Val Gln His Trp Pro Glu Thr Val Cys
 35 40 45
 5
 Glu Lys Ile Gln Asn Asp Cys Arg Asp Pro Pro Asp Tyr Trp Thr Ile
 50 55 60
 10
 His Gly Leu Trp Pro Asp Lys Ser Glu Gly Cys Asn Arg Ser Trp Pro
 65 70 75 80
 Phe Asn Leu Glu Glu Ile Lys Asp Leu Leu Pro Glu Met Arg Ala Tyr
 85 90 95
 15
 Trp Pro Asp Val Ile His Ser Phe Pro Asn Arg Ser Arg Phe Trp Lys
 100 105 110
 His Glu Trp Glu Lys His Gly Thr Cys Ala Ala Gln Val Asp Ala Leu
 115 120 125
 20
 Asn Ser Gln Lys Lys Tyr Phe Gly Arg Ser Leu Glu Leu Tyr Arg Glu
 130 135 140
 Leu Asp Leu Asn Ser Val Leu Leu Lys Leu Gly Ile Lys Pro Ser Ile
 145 150 155 160
 25
 Asn Tyr Tyr Gln Val Ala Asp Phe Lys Asp Ala Leu Ala Arg Val Tyr
 165 170 175
 30
 Gly Val Ile Pro Lys Ile Gln Cys Leu Pro Pro Ser Gln Asp Glu Glu
 180 185 190
 Val Gln Thr Ile Gly Gln Ile Glu Leu Cys Leu Thr Lys Gln Asp Gln
 195 200 205
 35
 Gln Leu Gln Asn Cys Thr Glu Pro Gly Glu Gln Pro Ser Pro Lys Gln
 210 215 220
 Glu Val Trp Leu Ala Asn Gly Ala Ala Glu Ser Arg Gly Leu Arg Val
 225 230 235 240
 40
 Cys Glu Asp Gly Pro Val Phe Tyr Pro Pro Pro Lys Lys Thr Lys His
 245 250 255
 45

50 (2) INFORMATION FOR SEQ ID NO: 181:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 324 amino acids:

(B) TYPE: amino acid

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 181:

Met Ala Pro Leu Leu Leu Gln Leu Ala Val Leu Gly Ala Ala Leu Ala
 1 5 10 15

60

309

Ala Ala Ala Leu Val Leu Ile Ser Ile Val Ala Phe Thr Thr Ala Thr
20 25 30

5 Lys Met Pro Ala Leu His Arg His Glu Glu Glu Lys Phe Phe Leu Asn
35 40 45

Ala Lys Gly Gln Lys Glu Thr Leu Pro Ser Ile Trp Asp Ser Pro Thr
50 55 60

10 Lys Gln Leu Ser Val Val Val Pro Ser Tyr Asn Glu Glu Lys Arg Leu
65 70 75 80

Pro Val Met Met Asp Glu Ala Leu Ser Tyr Leu Glu Lys Arg Gln Lys
85 90 95

15 Arg Asp Pro Ala Phe Thr Tyr Glu Val Ile Val Val Asp Asp Gly Ser
100 105 110

20 Lys Asp Gln Thr Ser Lys Val Ala Phe Lys Tyr Cys Gln Lys Tyr Gly
115 120 125

Ser Asp Lys Val Arg Val Ile Thr Leu Val Lys Asn Arg Gly Lys Gly
130 135 140

25 Gly Ala Ile Arg Met Gly Ile Phe Ser Ser Arg Gly Glu Lys Ile Leu
145 150 155 160

Met Ala Asp Ala Asp Gly Ala Thr Lys Phe Pro Asp Val Glu Lys Leu
165 170 175

30 Glu Lys Gly Leu Asn Asp Leu Gln Pro Trp Pro Asn Gln Met Ala Ile
180 185 190

35 Ala Cys Gly Ser Arg Ala His Leu Glu Lys Glu Ser Ile Ala Gln Arg
195 200 205

Ser Tyr Phe Arg Thr Leu Leu Met Tyr Gly Phe His Phe Leu Val Trp
210 215 220

40 Phe Leu Cys Val Lys Gly Ile Arg Asp Thr Gln Cys Gly Phe Lys Leu
225 230 235 240

Phe Thr Arg Glu Ala Ala Ser Arg Thr Phe Ser Ser Leu His Val Glu
245 250 255

45 Arg Trp Ala Phe Asp Val Glu Leu Leu Tyr Ile Ala Gln Phe Phe Lys
260 265 270

50 Ile Pro Ile Ala Glu Ile Ala Val Asn Trp Thr Glu Ile Glu Gly Ser
275 280 285

Lys Leu Val Pro Phe Trp Ser Trp Leu Gln Met Gly Lys Asp Leu Leu
290 295 300

55 Phe Ile Arg Leu Arg Tyr Leu Thr Gly Ala Trp Arg Leu Glu Gln Thr
305 310 315 320

Arg Lys Met Asn

60

(2) INFORMATION FOR SEQ ID NO: 182:

5

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 47 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 182:

10

Met Asp Ile Cys Phe Phe His Tyr Val Leu Leu Phe Phe Leu Val Arg
 1 5 10 15

15

Cys Ala Leu Val Val Leu Ile Leu Leu Cys Gln Gly Trp Gly Asn Gly
 20 25 30

Gly Gly Cys Val Gly Arg Val Leu Ile Ile Val Phe Ser Ser Val
 35 40 45

20

(2) INFORMATION FOR SEQ ID NO: 183:

25

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 93 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 183:

30

Met Ala Ser Leu Gly His Ile Leu Val Phe Cys Val Gly Leu Leu Thr
 1 5 10 15

Met Ala Lys Ala Glu Ser Pro Lys Glu His Asp Pro Phe Thr Tyr Asp
 20 25 30

35

Tyr Gln Ser Leu Gln Ile Gly Gly Leu Val Ile Ala Gly Ile Leu Phe
 35 40 45

40

Ile Leu Gly Ile Leu Ile Val Leu Ser Arg Arg Cys Arg Cys Lys Phe
 50 55 60

Asn Gln Gln Gln Arg Thr Gly Glu Pro Asp Glu Glu Glu Gly Thr Phe
 65 70 75 80

45

Arg Ser Ser Ile Arg Arg Leu Ser Thr Arg Arg Arg Xaa
 85 90

50

(2) INFORMATION FOR SEQ ID NO: 184:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 168 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 184:

Met Xaa Thr Lys Glu Phe Gly Xaa Gly Arg Ala Val Gln Gln Val Leu
 1 5 10 15

60

311

Asn Ile Glu Cys Leu Arg Asp Phe Leu Thr Pro Pro Leu Leu Ser Val
 20 25 30
 5 Arg Phe Arg Tyr Val Gly Ala Pro Gln Ala Leu Thr Leu Lys Leu Pro
 35 40 45
 Val Thr Xaa Asn Lys Phe Phe Gln Pro Thr Glu Met Ala Ala Gln Asp
 50 55 60
 10 Phe Phe Gln Arg Trp Lys Gln Leu Ser Leu Pro Gln Gln Glu Ala Gln
 65 70 75 80
 Lys Ile Phe Lys Ala Asn His Pro Met Asp Ala Glu Val Thr Lys Ala
 85 90 95
 15 Lys Leu Leu Gly Phe Gly Ser Ala Leu Leu Asp Asn Val Asp Pro Asn
 100 105 110
 20 Pro Glu Asn Phe Val Gly Ala Gly Ile Ile Gln Thr Lys Ala Leu Gln
 115 120 125
 Val Gly Cys Leu Leu Arg Leu Glu Pro Asn Ala Gln Ala Gln Met Tyr
 130 135 140
 25 Arg Leu Thr Leu Arg Thr Ser Lys Glu Pro Val Ser Arg His Leu Cys
 145 150 155 160
 Glu Leu Leu Ala Gln Gln Phe Xaa
 165
 30

(2) INFORMATION FOR SEQ ID NO: 185:

35 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 43 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 185:
 40 Met Phe Tyr Val Leu Ser Val Ser Pro Leu Leu Xaa Phe Leu Ala Cys
 1 5 10 15
 45 Gly Leu Cys Leu Cys Val Asn Trp Lys Ile Ala Ile Ser Gln Leu Ser
 20 25 30
 Leu Ser Phe Lys Asn Glu Leu Glu Lys Pro Xaa
 35 40
 50

(2) INFORMATION FOR SEQ ID NO: 186:

55 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 59 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 186:
 60 Met Lys Leu Phe Asp Ala Ser Pro Thr Phe Phe Ala Phe Leu Leu Gly

312

1 5 10 15
 His Ile Leu Ala Met Glu Val Leu Ala Trp Leu Leu Ile Tyr Leu Leu
 20 25 30
 5 Gly Pro Gly Trp Val Pro Ser Ala Leu Xaa Arg Leu His Pro Gly His
 35 40 45
 10 Leu Ser Gly Ser Val Leu Val Ser Ala Ala Xaa
 50 55

15 (2) INFORMATION FOR SEQ ID NO: 187:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 189 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 187:

Met Asp Val Asn Ile Ala Pro Leu Arg Ala Trp Asp Asp Phe Phe Pro
 1 5 10 15
 25 Gly Ser Asp Arg Phe Ala Arg Pro Asp Phe Arg Asp Ile Ser Lys Trp
 20 25 30
 Asn Asn Arg Val Val Ser Asn Leu Leu Tyr Tyr Gln Thr Asn Tyr Leu
 35 40 45
 30 Val Val Ala Ala Met Met Ile Ser Ile Val Gly Phe Leu Ser Pro Phe
 50 55 60
 35 Asn Met Ile Leu Gly Gly Ile Val Val Val Leu Val Phe Thr Gly Phe
 65 70 75 80
 Val Trp Ala Ala His Asn Lys Asp Val Leu Arg Arg Met Lys Lys Arg
 85 90 95
 40 Tyr Pro Thr Thr Phe Val Met Val Val Met Leu Ala Ser Tyr Phe Leu
 100 105 110
 Ile Ser Met Phe Gly Gly Val Met Val Phe Val Phe Gly Ile Thr Phe
 115 120 125
 45 Pro Leu Leu Leu Met Phe Ile His Ala Ser Leu Arg Leu Arg Asn Leu
 130 135 140
 Lys Asn Lys Leu Glu Asn Lys Met Glu Gly Ile Gly Leu Lys Arg Thr
 145 150 155 160
 Pro Met Gly Ile Val Leu Asp Ala Leu Glu Gln Gln Glu Glu Gly Ile
 165 170 175
 55 Asn Arg Leu Thr Asp Tyr Ile Ser Lys Val Lys Glu Xaa
 180 185

60 (2) INFORMATION FOR SEQ ID NO: 188:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 146 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 188:

5 Met Phe Leu Thr Arg Ile Leu Cys Pro Thr Tyr Ile Ala Leu Thr Phe
 1 5 10 15
 10 Leu Val Tyr Ile Val Ala Leu Val Ser Gly Gln Leu Cys Met Glu Ile
 20 25 30
 15 Ala Arg Gly Asn Ile Phe Phe Leu Asn Glu Leu Val Thr Thr Phe Cys
 35 40 45
 Cys Ser Cys Leu Leu Leu Ser Val Pro Tyr Leu His Pro Gly Phe Phe
 50 55 60
 20 Tyr Ser Ser Leu Cys Lys Cys Cys Phe Val Leu Val Val Leu Ser Arg
 65 70 75 80
 Ile Gly Ser Val Asn Glu Thr Trp Ser Cys Asn Phe Ser Ile Cys Ser
 85 90 95
 25 Tyr Leu Ile Phe Gly Ser Pro Ile Phe Thr Ala Val Ile Pro Lys Arg
 100 105 110
 30 Cys Ala Leu Glu Asp Ile Gln Asn Asn Pro Ile Gly Cys Leu Leu Arg
 115 120 125
 Cys Thr Pro Ala Trp Glu Thr Glu Gly Asp Ser Ile Ser Lys Lys Ile
 130 135 140
 35 Lys Lys
 145

(2) INFORMATION FOR SEQ ID NO: 189:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 84 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 189:

50 Met Gly Ser Arg Ala Glu Leu Cys Thr Leu Leu Gly Gly Phe Ser Phe
 1 5 10 15
 Leu Leu Leu Leu Ile Pro Gly Glu Gly Ala Lys Gly Gly Ser Leu Arg
 20 25 30
 55 Glu Ser Gln Gly Val Cys Ser Lys Gln Thr Leu Val Val Pro Leu His
 35 40 45
 Tyr Asn Glu Ser Tyr Ser Gln Pro Val Tyr Lys Pro Tyr Leu Thr Leu
 50 55 60
 60 Cys Ala Gly Ser Ala Ser Ala Ala Leu Thr Gly Pro Cys Thr Ala Leu

314

65 70 75 80

Cys Gly Gly Arg

5

(2) INFORMATION FOR SEQ ID NO: 190:

10

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 58 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 190:

15

Met Met Gly Val Leu Gln Leu Leu His Ile Phe Trp Ala Tyr Leu Ile
1 5 10 15

20

Leu Arg Met Ala His Lys Phe Ile Thr Gly Lys Leu Val Glu Asp Glu
20 25 30

Arg Ser Thr Gly Lys Lys Gln Arg Ala Gln Arg Gly Arg Arg Leu Gln
35 40 45

25

Leu Gly Glu Glu Gln Arg Ala Gly Pro Xaa
50 55

30

(2) INFORMATION FOR SEQ ID NO: 191:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 311 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 191:

35

Met Arg Arg Leu Val His Asp Leu Leu Pro Pro Glu Val Cys Ser Leu
1 5 10 15

40

Leu Asn Pro Ala Ala Ile Tyr Ala Asn Asn Glu Ile Ser Leu Arg Asp
20 25 30

45

Val Glu Val Tyr Gly Phe Asp Tyr Asp Tyr Thr Leu Ala Gln Tyr Ala
35 40 45

Asp Ala Leu His Pro Glu Ile Phe Ser Thr Ala Arg Asp Ile Leu Ile
50 55 60

50

Glu His Tyr Lys Tyr Pro Glu Gly Ile Arg Lys Tyr Asp Tyr Asn Pro
65 70 75 80

Ser Phe Ala Ile Arg Gly Leu His Tyr Asp Ile Gln Lys Ser Leu Leu
85 90 95

55

Met Lys Ile Asp Ala Phe His Tyr Val Gln Leu Gly Thr Ala Tyr Arg
100 105 110

60

Gly Leu Gln Pro Val Pro Asp Glu Glu Val Ile Glu Leu Tyr Gly Gly
115 120 125

315

Thr Gln His Ile Pro Leu Tyr Gln Met Ser Gly Phe Tyr Gly Lys Gly
 130 135 140

5 Pro Ser Ile Lys Gln Phe Met Asp Ile Phe Ser Leu Pro Glu Met Ala
 145 150 155 160

Leu Leu Ser Cys Val Val Asp Tyr Phe Leu Gly His Ser Leu Glu Phe
 165 170 175

10 Asp Gln Ala His Leu Tyr Lys Asp Val Thr Asp Ala Ile Arg Asp Val
 180 185 190

15 His Val Lys Gly Leu Met Tyr Gln Trp Ile Glu Gln Asp Met Glu Lys
 195 200 205

Tyr Ile Leu Arg Gly Asp Glu Thr Phe Ala Val Leu Ser Arg Leu Val
 210 215 220

20 Ala His Gly Lys Gln Leu Phe Leu Ile Thr Asn Ser Pro Phe Ser Phe
 225 230 235 240

Val Asp Lys Gly Met Arg His Met Val Gly Pro Asp Trp Arg His Ser
 245 250 255

25 Ser Met Trp Ser Leu Ser Arg Gln Thr Ser Pro Ala Ser Ser Leu Thr
 260 265 270

30 Gly Ala Ser Phe Xaa Glu Asn Ser Met Arg Arg Ala His Phe Ser Gly
 275 280 285

Thr Gly Ser Pro Ala Trp Lys Arg Ala Arg Ser Ile Gly Arg Glu Thr
 290 295 300

35 Cys Leu Thr Ser Tyr Ala Xaa
 305 310

40 (2) INFORMATION FOR SEQ ID NO: 192:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 318 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 192:

Met Asn Trp Glu Leu Leu Leu Trp Leu Leu Val Leu Cys Ala Leu Leu
 1 5 10 15

50 Leu Leu Leu Val Gln Leu Leu Arg Phe Leu Asn Ala Asp Gly Asp Leu
 20 25 30

Thr Leu Leu Trp Ala Glu Trp Gln Gly Arg Asn Pro Gln Trp Glu Leu
 35 40 45

Thr Asp Met Val Val Trp Val Thr Gly Ala Ser Leu Gly Ile Gly Glu
 50 55 60

60 Glu Leu Ala Tyr Gln Leu Ser Lys Leu Gly Thr Asn Leu Val Leu Ser

316

	65					70						75				80
	Ala	Arg	Arg	Val	His	Glu	Leu	Glu	Arg	Val	Lys	Arg	Arg	Cys	Leu	Glu
					85					90					95	
5	Asn	Gly	Asn	Leu	Lys	Glu	Lys	Asp	Ile	Leu	Val	Leu	Pro	Leu	Asp	Leu
				100					105					110		
	Thr	Asp	Thr	Gly	Ser	His	Glu	Ala	Ala	Thr	Lys	Ala	Val	Leu	Gln	Glu
10				115				120					125			
	Phe	Gly	Arg	Ile	Asp	Ile	Leu	Val	Asn	Asn	Gly	Gly	Met	Ser	Gln	Arg
		130					135					140				
15	Ser	Leu	Cys	Met	Asp	Thr	Ser	Leu	Asp	Val	Tyr	Arg	Lys	Leu	Ile	Glu
	145					150					155					160
	Leu	Asn	Tyr	Leu	Gly	Thr	Val	Ser	Leu	Thr	Lys	Cys	Val	Leu	Pro	His
20					165					170					175	
	Met	Ile	Glu	Arg	Lys	Gln	Gly	Lys	Ile	Val	Thr	Val	Asn	Ser	Ile	Leu
				180					185					190		
	Gly	Ile	Ile	Ser	Val	Pro	Leu	Ser	Ile	Gly	Tyr	Cys	Ala	Ser	Lys	His
25			195					200					205			
	Ala	Leu	Arg	Gly	Phe	Phe	Asn	Gly	Leu	Arg	Thr	Glu	Leu	Ala	Thr	Tyr
		210					215					220				
30	Pro	Gly	Ile	Ile	Val	Ser	Asn	Ile	Cys	Pro	Gly	Pro	Val	Gln	Ser	Asn
	225					230					235					240
	Ile	Val	Glu	Asn	Ser	Leu	Ala	Gly	Glu	Val	Thr	Lys	Thr	Ile	Gly	Asn
35					245					250					255	
	Asn	Gly	Asp	Gln	Ser	His	Lys	Met	Thr	Thr	Ser	Arg	Cys	Val	Arg	Leu
				260					265					270		
40	Met	Leu	Ile	Ser	Met	Ala	Asn	Asp	Leu	Lys	Glu	Val	Trp	Ile	Ser	Glu
		275						280					285			
	Gln	Pro	Phe	Leu	Phe	Ser	Asn	Ile	Phe	Val	Ala	Ile	His	Ala	Asn	Leu
		290					295					300				
45	Gly	Leu	Val	Asp	Asn	Gln	Gln	Asp	Gly	Glu	Glu	Lys	Asp	Xaa		
	305					310					315					

50 (2) INFORMATION FOR SEQ ID NO: 193:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 53 amino acids.

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ 11 3 100

Met Trp Pro Ser Phe Pro Gln Val Arg Val Leu Ile Phe Leu Phe Gly
1 5 10 15

60

317

Ile Leu Phe Phe Ser Phe Gly Ser Ser Ser Leu Pro Pro Gly Leu Pro
 20 25 30

Pro Pro Ala Ser Leu Leu Cys Cys Ala Val Gln Trp Gly Ala Arg Ala
 5 35 40 45

Leu Phe Leu Pro Ala
 50

10

(2) INFORMATION FOR SEQ ID NO: 194:

(i) SEQUENCE CHARACTERISTICS:
 15 (A) LENGTH: 42 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 194:

Met Leu Val Thr Cys Ser Val Cys Cys Tyr Leu Phe Trp Leu Ile Ala
 1 5 10 15

Ile Leu Ala Gln Leu Asn Pro Leu Phe Gly Pro Gln Leu Lys Asn Glu
 20 25 30

Thr Ile Trp Tyr Leu Lys Tyr His Trp Pro
 35 40

30

(2) INFORMATION FOR SEQ ID NO: 195:

(i) SEQUENCE CHARACTERISTICS:
 35 (A) LENGTH: 102 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 195:

Met Glu Gly Thr Glu Met Gly Ala Arg Pro Gly Gly His Pro Gln Lys
 40 1 5 10 15

Trp Ser Phe Leu Trp Ser Leu Ala Leu Trp Leu Pro Leu Ala Leu Ser
 20 25 30

Val Ser Leu Phe Leu Gly Leu Ser Leu Ser Pro Pro Gln Pro Gly Leu
 45 35 40 45

Ser Leu Trp Cys Thr Leu Ser Tyr Cys Cys Glu Gln Trp Lys Phe Lys
 50 55 60

Gly Thr Pro Ser Pro Ala Leu Leu Asn Leu Gly Thr Gln Pro Lys Lys
 65 70 75 80

Asp Lys Lys Leu Glu Asp Ser Ile Ala Thr Gln Leu Arg Glu Leu Pro
 55 85 90 95

Glu Lys Asn Ser Asn Xaa
 100

60

(2) INFORMATION FOR SEQ ID NO: 196:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 45 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 196:

10 Met Ala Leu Thr Phe Leu Leu Val Leu Leu Thr Leu Ala Thr Ser Ala
1 5 10 15
His Gly Cys Thr Glu Thr Ser Asp Ala Gly Arg Ala Ser Thr Gly Gly
20 25 30
15 Pro Gln Arg Thr Ala Arg Thr Gln Trp Leu Leu Cys Xaa
35 40 45

20

(2) INFORMATION FOR SEQ ID NO: 197:

(i) SEQUENCE CHARACTERISTICS:

- 25 (A) LENGTH: 355 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 197:

30 Met Gly Pro Ser Thr Pro Leu Leu Ile Leu Phe Leu Leu Ser Trp Ser
1 5 10 15
Gly Pro Leu Gln Gly Gln Gln His His Leu Val Glu Tyr Met Glu Arg
20 25 30
35 Arg Leu Ala Ala Leu Glu Glu Arg Leu Ala Gln Cys Gln Asp Gln Ser
35 40 45
Ser Arg His Ala Ala Glu Leu Arg Asp Phe Lys Asn Lys Met Leu Pro
50 55 60
40 Leu Leu Glu Val Ala Glu Lys Glu Arg Glu Ala Leu Arg Thr Glu Ala
65 70 75 80
45 Asp Thr Ile Ser Gly Arg Val Asp Arg Leu Glu Arg Glu Val Asp Tyr
85 90 95
Leu Glu Thr Gln Asn Pro Ala Leu Pro Cys Val Glu Phe Asp Glu Lys
100 105 110
50 Val Thr Gly Gly Pro Gly Thr Lys Gly Lys Gly Arg Arg Asn Glu Lys
115 120 125
Tyr Asp Met Val Thr Asp Cys Gly Tyr Thr Ile Ser Gln Val Arg Ser
130 135 140
55 Met Lys Ile Leu Lys Arg Phe Gly Gly Pro Ala Gly Leu Trp Thr Lys
145 150 155 160
60 Asp Pro Leu Gly Gln Thr Glu Lys Ile Tyr Val Leu Asp Gly Thr Gln
165 170 175

319

Asn Asp Thr Ala Phe Val Phe Pro Arg Leu Arg Asp Phe Thr Leu Ala
 180 185 190
 5 Met Ala Ala Arg Lys Ala Ser Arg Val Arg Val Pro Phe Pro Trp Val
 195 200 205
 Gly Thr Gly Gln Leu Val Tyr Gly Gly Phe Leu Tyr Phe Ala Arg Arg
 210 215 220
 10 Pro Pro Gly Arg Pro Gly Gly Gly Gly Glu Met Glu Asn Thr Leu Gln
 225 230 235 240
 Leu Ile Lys Phe His Leu Ala Asn Arg Thr Val Val Asp Ser Ser Val
 15 245 250 255
 Phe Pro Ala Glu Gly Leu Ile Pro Pro Tyr Gly Leu Thr Ala Asp Thr
 260 265 270
 20 Tyr Ile Asp Leu Ala Ala Asp Glu Glu Gly Leu Trp Ala Val Tyr Ala
 275 280 285
 Thr Arg Glu Asp Asp Arg His Leu Cys Leu Ala Lys Leu Asp Pro Gln
 290 295 300
 25 Thr Leu Asp Thr Glu Gln Gln Trp Asp Thr Pro Cys Pro Arg Glu Asn
 305 310 315 320
 Ala Glu Ala Ala Phe Val Ile Cys Gly Thr Leu Tyr Val Val Tyr Asn
 30 325 330 335
 Thr Arg Pro Ala Ser Arg Ala Arg Ile Gln Cys Ser Phe Asp Ala Ser
 340 345 350
 35 Gly Pro Xaa
 355

40 (2) INFORMATION FOR SEQ ID NO: 198:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 74 amino acids

(B) TYPE: amino acid

45 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 198:

Met Val Leu Pro Leu Leu Ile Phe Val Leu Leu Pro Lys Val Val Asn
 1 5 10 15
 50 Thr Ser Asp Pro Asp Met Arg Arg Glu Met Glu Gln Ser Met Asn Met
 20 25 30
 Leu Asn Ser Asn His Glu Leu Pro Asp Val Ser Glu Phe Met Thr Arg
 35 40 45
 55 Leu Phe Ser Ser Lys Ser Ser Gly Lys Ser Ser Ser Gly Ser Ser Lys
 50 55 60
 60 Thr Gly Lys Ser Gly Ala Gly Lys Arg Arg

320

65

70

5 (2) INFORMATION FOR SEQ ID NO: 199:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 113 amino acids

(B) TYPE: amino acid

10 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 199:

Met Phe Thr Met Leu Cys Ile Asn Gly Thr Thr Pro Arg Pro Leu Pro
 1 5 10 15
 Val Pro Ser Pro Phe Gly Cys Met Ile Phe Phe Phe Phe Lys Asn Pro
 20 25 30
 Trp Lys Gln Arg Leu Leu Gln Gly Trp Leu Gly Ala Arg Pro Ile His
 35 40 45
 Leu Leu Gly Tyr Leu Pro Leu Ser Leu Leu Trp Cys Pro Phe Pro Leu
 50 55 60
 Pro Cys Ala Arg Cys Ser Val Val Tyr Ile Ser Ser Pro Arg His Gly
 65 70 75 80
 Ala His Ala Pro Arg Asp Met Ile Leu Ser Leu Val Leu Ala His Gly
 85 90 95
 Ala Leu Tyr Lys Glu Leu Gly Gly Arg Gly Arg Lys Trp Glu Pro Ser
 100 105 110
 Xaa

35

40 (2) INFORMATION FOR SEQ ID NO: 200:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 123 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 200:

Met Ala Cys Arg Cys Leu Ser Phe Leu Leu Met Gly Thr Phe Leu Ser
 1 5 10 15
 Val Ser Gln Thr Val Leu Ala Gln Leu Asp Ala Leu Leu Val Phe Pro
 20 25 30
 Gly Gln Val Ala Gln Leu Ser Cys Thr Leu Ser Pro Gln His Val Thr
 35 40 45
 Ile Arg Asp Tyr Gly Val Ser Trp Tyr Gln Gln Arg Ala Gly Ser Ala
 50 55 60
 Pro Arg Tyr Leu Leu Tyr Tyr Arg Ser Glu Glu Asp His His Arg Pro
 65 70 75 80

60

321

Ala Asp Ile Pro Asp Arg Phe Ser Ala Ala Lys Asp Glu Ala His Asn
85 90 95

5 Ala Cys Val Leu Thr Ile Ser Pro Val Gln Pro Glu Asp Asp Ala Asp
100 105 110

Tyr Tyr Cys Ser Val Gly Tyr Gly Phe Ser Pro
115 120

10

(2) INFORMATION FOR SEQ ID NO: 201:

15 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 315 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 201:

20 Met Ala Gly Gly Arg Cys Gly Pro Xaa Leu Thr Ala Leu Leu Ala Ala
1 5 10 15

25 Trp Ile Ala Ala Val Ala Ala Thr Ala Gly Pro Glu Glu Ala Ala Leu
20 25 30

Pro Pro Glu Gln Ser Arg Val Gln Pro Met Thr Ala Ser Asn Trp Thr
35 40 45

30 Leu Val Met Glu Gly Glu Trp Met Leu Lys Phe Tyr Ala Pro Trp Cys
50 55 60

Pro Ser Cys Gln Gln Thr Asp Ser Glu Trp Glu Ala Phe Ala Lys Asn
65 70 75 80

35 Gly Glu Ile Leu Gln Ile Ser Val Gly Lys Val Asp Val Ile Gln Glu
85 90 95

40 Pro Gly Leu Ser Gly Arg Phe Phe Val Thr Thr Leu Pro Ala Phe Phe
100 105 110

His Ala Lys Asp Gly Ile Phe Arg Arg Tyr Arg Gly Pro Gly Ile Phe
115 120 125

45 Glu Asp Leu Gln Asn Tyr Ile Leu Glu Lys Lys Trp Gln Ser Val Glu
130 135 140

Pro Leu Thr Gly Trp Lys Ser Pro Ala Ser Leu Thr Met Ser Gly Met
145 150 155 160

50 Ala Gly Leu Phe Ser Ile Ser Gly Lys Ile Trp His Leu His Asn Tyr
165 170 175

55 Phe Thr Val Thr Leu Gly Ile Pro Ala Trp Cys Ser Tyr Val Phe Phe
180 185 190

Val Ile Ala Thr Leu Val Phe Gly Leu Phe Met Gly Leu Val Leu Val
195 200 205

60 Val Ile Ser Glu Cys Phe Tyr Val Pro Leu Pro Arg His Leu Ser Glu

322

210 215 220
 Arg Ser Glu Gln Asn Arg Arg Ser Glu Glu Ala His Arg Ala Glu Gln
 225 230 235 240
 5 Leu Gln Asp Ala Glu Glu Glu Lys Asp Asp Ser Asn Glu Glu Glu Asn
 245 250 255
 Lys Asp Ser Leu Val Asp Asp Glu Glu Glu Lys Glu Asp Leu Gly Asp
 10 260 265 270
 Glu Asp Glu Ala Glu Glu Glu Glu Glu Glu Asp Asn Leu Ala Ala Gly
 275 280 285
 15 Val Asp Glu Glu Arg Ser Glu Ala Asn Asp Gln Gly Pro Pro Gly Glu
 290 295 300
 Asp Gly Val Thr Arg Glu Xaa Ser Arg Ala Xaa
 20 305 310 315
 (2) INFORMATION FOR SEQ ID NO: 202:
 25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 236 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 202:
 30 Met Gly Thr Ala Asp Ser Asp Glu Met Ala Pro Glu Ala Pro Gln His
 1 5 10 15
 Thr His Ile Asp Val His Ile His Gln Glu Ser Ala Leu Ala Lys Leu
 35 20 25 30
 Leu Leu Thr Cys Cys Ser Ala Leu Arg Pro Arg Ala Thr Gln Ala Arg
 35 40 45
 40 Gly Ser Ser Arg Leu Leu Val Ala Ser Trp Val Met Gln Ile Val Leu
 50 55 60
 Gly Ile Leu Ser Ala Val Leu Gly Gly Phe Phe Tyr Ile Arg Asp Tyr
 45 65 70 75 80
 Thr Leu Leu Val Thr Ser Gly Ala Ala Ile Trp Thr Gly Ala Val Ala
 85 90 95
 Val Leu Ala Gly Ala Ala Ala Phe Ile Tyr Glu Lys Arg Gly Gly Thr
 50 100 105 110
 Tyr Trp Ala Leu Leu Arg Thr Leu Leu Ala Leu Ala Ala Phe Ser Thr
 115 120 125
 55 Ala Ile Ala Ala Leu Lys Leu Trp Asn Glu Asp Phe Arg Tyr Gly Tyr
 130 135 140
 Ser Tyr Tyr Asn Ser Ala Cys Arg Ile Ser Thr Ser Ser Asp Trp Asn
 60 145 150 155 160

323

Thr Pro Ala Pro Thr Gln Ser Pro Glu Glu Val Arg Arg Leu His Leu
 165 170 175
 5 Cys Thr Ser Phe Met Asp Met Leu Lys Ala Leu Phe Arg Thr Leu Gln
 180 185 190
 Ala Met Leu Leu Gly Val Trp Ile Leu Leu Leu Leu Ala Ser Leu Ala
 195 200 205
 10 Pro Leu Trp Leu Tyr Cys Trp Arg Met Phe Pro Thr Lys Gly Lys Arg
 210 215 220
 Asp Gln Lys Glu Met Leu Glu Val Ser Gly Ile Xaa
 225 230 235
 15

(2) INFORMATION FOR SEQ ID NO: 203:

20 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 93 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 203:

25 Met Ile His Leu Gly His Ile Leu Phe Leu Leu Leu Leu Pro Val Ala
 1 5 10 15
 30 Ala Ala Gln Thr Thr Pro Gly Glu Arg Ser Ser Leu Pro Ala Phe Tyr
 20 25 30
 Pro Gly Thr Ser Gly Ser Cys Ser Gly Cys Gly Ser Leu Ser Leu Pro
 35 40 45
 35 Leu Leu Ala Gly Leu Val Ala Ala Asp Ala Val Ala Ser Leu Leu Ile
 50 55 60
 Val Gly Ala Val Phe Leu Cys Ala Arg Pro Arg Arg Ser Pro Ala Gln
 65 70 75 80
 40 Glu Asp Gly Lys Val Tyr Ile Asn Met Pro Gly Arg Gly
 85 90

45

(2) INFORMATION FOR SEQ ID NO: 204:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 35 amino acids
 50 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 204:

55 Met Trp Ser Ala Gly Arg Gly Gly Ala Ala Trp Pro Val Leu Leu Gly
 1 5 10 15
 Leu Leu Leu Ala Leu Leu Val Pro Gly Gly Gly Ala Ala Lys Thr Gly
 20 25 30
 60 Ala Asp Ser

35

5 (2) INFORMATION FOR SEQ ID NO: 205:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 43 amino acids

(B) TYPE: amino acid

10 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 205:

Asp Cys Xaa His Val Ser Val Leu Gln Ser Thr Ile Ser Pro Leu Leu
 1 5 10 15
 Pro Leu Pro Leu Leu Leu Pro His Gly Asn Cys Glu Glu Ala Pro Trp
 20 25 30
 Gln Ala Ala Val Ile Gly Gly Gly Asp Arg Ile
 35 40

25 (2) INFORMATION FOR SEQ ID NO: 206:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 85 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 206:

Met Arg Asp Cys Leu Ser Leu Lys Pro Arg Pro Leu Phe Pro Thr Gln
 1 5 10 15
 Phe Phe Phe Ile Leu Leu Leu Ile Phe Ile Ala Glu Val Ala Ala Ala
 20 25 30
 Val Val Ala Leu Val Tyr Thr Thr Met Val Arg His Trp Asp Gly Gly
 35 40 45
 Arg Glu Glu Asp Trp Ala Lys Pro Trp Glu Trp Ala Val Ala Cys Glu
 50 55 60
 Trp Pro Pro Ser Val Pro Ala Pro Lys His Trp Pro Ala Ser Pro Arg
 65 70 75 80
 Leu Ser Thr Ser Xaa
 85

50

(2) INFORMATION FOR SEQ ID NO: 207:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 208 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 207:

60 Met His Gly Asn Glu Ala Leu Gly Arg Glu Leu Leu Leu Leu Met

325

1 5 10 15
 Gln Phe Leu Cys His Glu Phe Leu Arg Xaa Asn Pro Arg Val Thr Arg
 20 25 30
 5
 Leu Leu Ser Glu Met Arg Ile His Leu Leu Pro Ser Met Asn Pro Asp
 35 40 45
 10
 Gly Tyr Glu Ile Ala Tyr His Arg Gly Ser Glu Leu Val Gly Trp Ala
 50 55 60
 Glu Gly Arg Trp Asn Asn Gln Ser Ile Asp Leu Asn His Asn Phe Ala
 65 70 75 80
 15
 Xaa Leu Asn Thr Pro Leu Trp Glu Ala Gln Asp Asp Gly Lys Val Pro
 85 90 95
 20
 His Ile Val Pro Asn His His Leu Pro Leu Pro Thr Tyr Tyr Thr Leu
 100 105 110
 Pro Asn Ala Thr Val Ala Pro Glu Thr Arg Ala Val Ile Lys Trp Met
 115 120 125
 25
 Lys Arg Ile Pro Phe Val Leu Ser Ala Asn Leu His Gly Gly Glu Leu
 130 135 140
 Val Val Ser Tyr Pro Phe Asp Met Thr Arg Thr Pro Trp Ala Ala Arg
 145 150 155 160
 30
 Glu Leu Thr Pro Thr Pro Asp Asp Ala Val Phe Arg Trp Leu Ser Thr
 165 170 175
 35
 Val Tyr Ala Gly Ser Asn Leu Ala Met Gln Asp Thr Ser Arg Arg Pro
 180 185 190
 Cys His Ser Gln Asp Phe Ser Val His Gly Asn Ile Ile Asn Gly Ala
 195 200 205
 40
 45
 (2) INFORMATION FOR SEQ ID NO: 208:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 24 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 50
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 208:
 Met Glu Ile Ser Cys Leu Leu Leu Leu Ile Gln Asp Ser Asp Glu Met
 1 5 10 15
 55
 Glu Asp Gly Pro Gly Val Gln Asp
 20
 60
 (2) INFORMATION FOR SEQ ID NO: 209:

326

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 483 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 209:

Met Ala Thr Gly Gly Gly Ile Arg Ala Met Thr Ser Leu Tyr Gly Gln
 1 5 10 15
 Leu Ala Gly Leu Lys Glu Leu Gly Leu Leu Asp Cys Xaa Ser Tyr Ile
 20 25 30
 Thr Gly Ala Ser Gly Ser Thr Trp Ala Leu Ala Asn Leu Tyr Lys Asp
 35 40 45
 Pro Glu Trp Ser Gln Lys Asp Leu Ala Gly Pro Thr Glu Leu Leu Lys
 50 55 60
 Thr Gln Val Thr Lys Asn Lys Leu Gly Val Leu Ala Pro Ser Gln Leu
 65 70 75 80
 Gln Arg Tyr Arg Gln Glu Leu Ala Glu Arg Ala Arg Leu Gly Tyr Pro
 85 90 95
 Ser Cys Phe Thr Asn Leu Trp Ala Leu Ile Asn Glu Ala Leu Leu His
 100 105 110
 Asp Glu Pro His Asp His Lys Leu Ser Asp Gln Arg Glu Ala Leu Ser
 115 120 125
 His Gly Gln Asn Pro Leu Pro Ile Tyr Cys Ala Leu Asn Thr Lys Gly
 130 135 140
 Gln Ser Leu Thr Thr Phe Glu Phe Gly Glu Trp Cys Glu Phe Ser Pro
 145 150 155 160
 Tyr Glu Val Gly Phe Pro Lys Tyr Gly Ala Phe Ile Pro Ser Glu Leu
 165 170 175
 Phe Gly Ser Glu Phe Phe Met Gly Gln Leu Met Lys Arg Leu Pro Glu
 180 185 190
 Ser Arg Ile Cys Phe Leu Glu Gly Ile Trp Ser Asn Leu Tyr Ala Ala
 195 200 205
 Asn Leu Gln Asp Ser Leu Tyr Trp Ala Ser Glu Pro Ser Gln Phe Trp
 210 215 220
 Asp Arg Trp Val Arg Asn Gln Ala Asn Leu Asp Lys Glu Gln Val Pro
 225 230 235 240
 Leu Leu Lys Ile Glu Glu Pro Pro Ser Thr Ala Gly Asn Ile Ala Glu
 245 250 255
 Phe Phe Thr Asp Leu Leu Thr Trp Arg Pro Leu Ala His Ala Thr His
 260 265 270
 Asn Phe Leu Arg Gly Leu His Phe His Lys Arg Tyr Phe Gln His Pro
 275 280 285

327

His Phe Ser Thr Trp Lys Ala Thr Thr Leu Asp Gly Leu Pro Asn Gln
 290 295 300
 5 Leu Thr Pro Ser Glu Pro His Leu Cys Leu Leu Asp Val Gly Tyr Leu
 305 310 315 320
 Ile Asn Thr Ser Cys Leu Pro Leu Leu Gln Pro Thr Arg Asp Val Asp
 325 330 335
 10 Leu Ile Leu Ser Leu Asp Tyr Asn Leu His Gly Ala Phe Gln Gln Leu
 340 345 350
 Gln Leu Leu Gly Arg Phe Cys Gln Glu Gln Gly Ile Pro Phe Pro Pro
 15 355 360 365
 Ile Ser Pro Ser Pro Glu Glu Gln Leu Gln Pro Arg Glu Cys His Thr
 370 375 380
 20 Phe Ser Asp Pro Thr Cys Pro Gly Ala Pro Ala Val Leu His Phe Pro
 385 390 395 400
 Leu Val Ser Asp Ser Phe Arg Glu Tyr Ser Ala Pro Gly Val Arg Arg
 405 410 415
 25 Thr Pro Glu Glu Ala Ala Ala Gly Glu Val Asn Leu Ser Ser Ser Asp
 420 425 430
 Ser Pro Tyr His Tyr Thr Lys Val Thr Tyr Ser Gln Glu Asp Val Asp
 10 435 440 445
 Lys Leu Leu His Leu Thr His Tyr Asn Val Cys Asn Asn Gln Glu Gln
 450 455 460
 35 Leu Leu Glu Ala Leu Arg Gln Ala Val Gln Arg Arg Arg Gln Arg Arg
 465 470 475 480
 Pro His Xaa

40

(2) INFORMATION FOR SEQ ID NO: 210:

45

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 13 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

50

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 210:

Leu Glu Val Gly Cys Ile Gln Val Ala Pro Asp Thr Phe
 1 5 10

55

(2) INFORMATION FOR SEQ ID NO: 211:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 20 amino acids
 (B) TYPE: amino acid

60

328

(D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 211:

5 Met Ser Leu Phe Phe Leu Leu Thr Leu Ile Ser Lys Leu His Gly Asp
 1 5 10 15
 Ala Glu Val Cys
 20

10

(2) INFORMATION FOR SEQ ID NO: 212:

15 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 55 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 212:

20 Met Pro His Pro Pro Leu Pro Glu Thr Ser Leu Glu Ala Gln Leu Pro
 1 5 10 15
 Met Gly Leu Leu Gln Leu Leu Arg Cys Ser Val Gln Ala Trp Ser Pro
 20 25 30
 25 Pro Pro Ser Ser Phe Cys Pro Gly Ser Glu Pro Arg Ser Ala Ser Ala
 35 40 45
 His Trp Gly Tyr Trp Trp Pro
 30 50 55

35 (2) INFORMATION FOR SEQ ID NO: 213:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 35 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 213:

Asp Pro Glu Thr Arg Trp His His Gly Gly Ser Ala Gln Asn Gly Leu
 1 5 10 15
 45 Leu Met Leu Ile Ser Val Leu Gln Gln Pro Val Ile Gly Thr Gly Ser
 20 25 30
 Tyr Leu Cys
 35

50

(2) INFORMATION FOR SEQ ID NO: 214:

55 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 230 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 214:

60

329

Met Glu Pro Leu Arg Leu Leu Ile Leu Leu Phe Val Thr Glu Leu Ser
 1 5 10 15
 Gly Ala His Asn Thr Thr Val Phe Gln Gly Val Ala Gly Gln Ser Leu
 5 20 25 30
 Gln Val Ser Cys Pro Tyr Asp Ser Met Lys His Trp Gly Arg Arg Lys
 35 40 45
 Ala Trp Cys Arg Gln Leu Gly Glu Lys Gly Pro Cys Gln Arg Val Val
 10 50 55 60
 Ser Thr His Asn Leu Trp Leu Leu Ser Phe Leu Arg Arg Trp Asn Gly
 15 65 70 75 80
 Ser Thr Ala Ile Thr Asp Asp Thr Leu Gly Gly Thr Leu Thr Ile Thr
 85 90 95
 Leu Arg Asn Leu Gln Pro His Asp Ala Gly Leu Tyr Gln Cys Gln Ser
 20 100 105 110
 Leu His Gly Ser Glu Ala Asp Thr Leu Arg Lys Val Leu Val Glu Val
 115 120 125
 Leu Ala Asp Pro Leu Asp His Arg Asp Ala Gly Asp Leu Trp Phe Pro
 25 130 135 140
 Gly Glu Ser Glu Ser Phe Glu Asp Ala His Val Glu His Ser Ile Ser
 145 150 155 160
 Arg Ser Leu Leu Glu Gly Glu Ile Pro Phe Pro Pro Thr Ser Ile Leu
 30 165 170 175
 Leu Leu Leu Ala Cys Ile Phe Leu Ile Lys Ile Leu Ala Ala Ser Xaa
 35 180 185 190
 Leu Trp Ala Ala Ala Trp His Gly Gln Lys Pro Gly Thr His Pro Pro
 195 200 205
 Ser Glu Leu Asp Cys Gly His Asp Pro Gly Tyr Gln Leu Gln Thr Leu
 40 210 215 220
 Pro Gly Leu Arg Asp Thr
 45 225 230

(2) INFORMATION FOR SEQ ID NO: 215:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 231 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 215:

55

Met Glu Pro Leu Arg Leu Leu Ile Leu Leu Phe Val Thr Glu Leu Ser
 1 5 10 15

60

Gly Ala His Asn Thr Thr Val Phe Gln Gly Val Ala Gly Gln Ser Leu
 20 25 30

330

Gln Val Ser Cys Pro Tyr Asp Ser Met Lys His Trp Gly Arg Arg Lys
 35 40 45
 5 Ala Trp Cys Arg Gln Leu Gly Glu Lys Gly Pro Cys Gln Arg Val Val
 50 55 60
 Ser Thr His Asn Leu Trp Leu Leu Ser Phe Leu Arg Arg Trp Asn Gly
 65 70 75 80
 10 Ser Thr Ala Ile Thr Asp Asp Thr Leu Gly Gly Thr Leu Thr Ile Thr
 85 90 95
 Leu Arg Asn Leu Gln Pro His Asp Ala Gly Leu Tyr Gln Cys Gln Ser
 15 100 105 110
 Leu His Gly Ser Glu Ala Asp Thr Leu Arg Lys Val Leu Val Glu Val
 115 120 125
 20 Leu Ala Asp Pro Leu Asp His Arg Asp Ala Gly Asp Leu Trp Phe Pro
 130 135 140
 Gly Glu Ser Glu Ser Phe Glu Asp Ala His Val Glu His Ser Ile Ser
 145 150 155 160
 25 Arg Ser Leu Leu Glu Gly Glu Ile Pro Phe Pro Pro Thr Ser Ile Leu
 165 170 175
 Leu Leu Leu Ala Cys Ile Phe Leu Ile Lys Ile Leu Ala Ala Ser Ala
 30 180 185 190
 Leu Trp Ala Ala Ala Trp His Gly Gln Lys Pro Gly Thr His Pro Pro
 195 200 205
 35 Ser Glu Leu Asp Cys Gly His Asp Pro Gly Tyr Gln Leu Gln Thr Leu
 210 215 220
 Pro Gly Leu Arg Asp Thr Xaa
 225 230
 40

(2) INFORMATION FOR SEQ ID NO: 216:

45 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 127 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 216:
 50 Met Gly Leu Thr Gly Phe Gly Val Phe Phe Leu Phe Phe Gly Met Ile
 1 5 10 15
 Leu Phe Phe Asp Lys Ala Leu Leu Ala Ile Gly Asn Val Leu Phe Val
 55 20 25 30
 Ala Gly Leu Ala Phe Val Ile Gly Leu Glu Arg Thr Phe Arg Phe Phe
 35 40 45
 60 Phe Gln Lys His Lys Met Lys Ala Thr Gly Phe Phe Leu Gly Gly Val

331

50 55 60

Phe Val Val Leu Ile Gly Trp Pro Leu Ile Gly Met Ile Phe Glu Ile
65 70 75 80

5 Tyr Gly Phe Phe Leu Leu Phe Arg Gly Phe Phe Pro Val Val Val Gly
85 90 95

10 Phe Ile Arg Arg Val Pro Val Leu Gly Ser Leu Leu Asn Leu Pro Gly
100 105 110

Ile Arg Ser Phe Val Asp Lys Val Gly Glu Ser Asn Asn Met Val
115 120 125

15

(2) INFORMATION FOR SEQ ID NO: 217:

(i) SEQUENCE CHARACTERISTICS:

20 (A) LENGTH: 47 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 217:

25 Met Ile Arg Lys Leu His Lys Ile Ile Val Phe Ser Pro Arg Val Ile
1 5 10 15

Val Leu Leu Asn Cys Phe Phe Phe Ile Lys Ala Lys Phe Val Leu Tyr
20 25 30

30 Ile Phe Val Phe His Val Leu Asp Gly Ser Ile Ser Tyr Pro Val
35 40 45

35

(2) INFORMATION FOR SEQ ID NO: 218:

(i) SEQUENCE CHARACTERISTICS:

40 (A) LENGTH: 41 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 218:

45 Met Leu Leu Asn Gln His Phe Lys Ile Phe Gly Ser Leu Ile His Met
1 5 10 15

Asn Leu Leu Phe Ala Leu Ile Ser Leu Gly Ser Ser Asn Leu Ser Gly
20 25 30

50 Val Gln Phe Cys Cys Glu Thr Val Gln
35 40

55 (2) INFORMATION FOR SEQ ID NO: 219:

(i) SEQUENCE CHARACTERISTICS:

60 (A) LENGTH: 105 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

332

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 219:

5 Met Gln Pro Leu Asn Phe Ser Ser Thr Xaa Cys Ser Ser Phe Ser Pro
 1 5 10 15
 Pro Thr Thr Val Ile Leu Leu Ile Leu Leu Cys Phe Glu Gly Leu Leu
 20 25 30
 10 Phe Leu Ile Phe Thr Ser Val Met Phe Gly Thr Gln Val His Ser Ile
 35 40 45
 Cys Thr Asp Glu Thr Gly Ile Glu Gln Leu Lys Lys Glu Glu Arg Arg
 50 55 60
 15 Trp Ala Lys Lys Thr Lys Trp Met Asn Met Lys Ala Val Phe Gly His
 65 70 75 80
 Pro Phe Ser Leu Gly Trp Ala Ser Pro Phe Ala Thr Pro Asp Gln Gly
 85 90 95
 20 Lys Ala Asp Pro Tyr Gln Tyr Val Val
 100 105

25

(2) INFORMATION FOR SEQ ID NO: 220:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 29 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 220:

35 Met Tyr Thr Asn His Phe Asn Leu Tyr Leu Lys Tyr Ile Leu Leu Ile
 1 5 10 15
 Ile Leu Ile Leu Asn Met Thr Asn Ser Ser Ser Arg Tyr
 20 25

40

(2) INFORMATION FOR SEQ ID NO: 221:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 17 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 221:

50 Met Asn Glu Leu Leu Leu Phe Phe Phe Phe Phe Phe Phe Leu His Phe
 1 5 10 15
 Val

55

(2) INFORMATION FOR SEQ ID NO: 222:

60 (i) SEQUENCE CHARACTERISTICS:

333

(A) LENGTH: 138 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 222:

5
 Met Lys Phe Thr Thr Leu Leu Phe Leu Ala Ala Val Ala Gly Ala Leu
 1 5 10 15
 10
 Val Tyr Ala Glu Asp Ala Ser Ser Asp Ser Thr Gly Ala Asp Pro Ala
 20 25 30
 Gln Glu Ala Gly Thr Ser Lys Pro Asn Glu Glu Ile Ser Gly Pro Ala
 35 40 45
 15
 Glu Pro Ala Ser Pro Pro Glu Thr Thr Thr Thr Ala Gln Glu Xaa Ser
 50 55 60
 Ala Ala Ala Val Gln Gly Thr Ala Lys Val Thr Ser Ser Arg Gln Glu
 65 70 75 80
 20
 Leu Asn Pro Leu Lys Ser Ile Val Glu Lys Ser Ile Leu Leu Thr Glu
 85 90 95
 Gln Ala Leu Ala Lys Ala Gly Lys Gly Met His Gly Gly Val Pro Gly
 100 105 110
 25
 Gly Lys Gln Phe Ile Glu Asn Gly Ser Glu Phe Ala Gln Lys Leu Leu
 115 120 125
 30
 Lys Lys Phe Ser Leu Leu Lys Pro Trp Ala
 130 135

35 (2) INFORMATION FOR SEQ ID NO: 223:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 50 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 223:

45
 Met Leu Gly Cys Gly Ile Pro Ala Leu Gly Leu Leu Leu Leu Gln
 1 5 10 15
 Xaa Ser Ala Asp Gly Asn Gly Ile Gln Gly Phe Phe Tyr Pro Trp Ser
 20 25 30
 50
 Cys Glu Gly Asp Ile Trp Asp Arg Glu Ser Cys Gly Gly Gln Ala Ala
 35 40 45
 Ile Arg
 50

55

(2) INFORMATION FOR SEQ ID NO: 224:

(i) SEQUENCE CHARACTERISTICS:

60 (A) LENGTH: 15 amino acids

334

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 224:

5 Met Glu Ala Val Phe Thr Val Phe Phe Phe Leu Leu Phe Cys Phe
 1 5 10 15

10 (2) INFORMATION FOR SEQ ID NO: 225:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 155 amino acids

(B) TYPE: amino acid

15 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 225:

Met Gly Phe Gly Ala Thr Leu Ala Val Gly Leu Thr Ile Phe Val Leu
 1 5 10 15

20 Ser Val Val Thr Ile Ile Ile Cys Phe Thr Cys Ser Cys Cys Cys Leu
 20 25 30

25 Tyr Lys Thr Cys Arg Arg Pro Arg Pro Val Val Thr Thr Thr Thr Ser
 35 40 45

Thr Thr Val Val His Ala Pro Tyr Pro Gln Pro Pro Ser Val Pro Pro
 50 55 60

30 Ser Tyr Pro Gly Pro Ser Tyr Gln Gly Tyr His Thr Met Pro Pro Gln
 65 70 75 80

Pro Gly Met Pro Ala Ala Pro Tyr Pro Met Gln Tyr Pro Pro Pro Tyr
 85 90 95

35 Pro Ala Gln Pro Met Gly Pro Pro Ala Tyr His Glu Thr Leu Ala Gly
 100 105 110

40 Gly Ala Ala Ala Pro Tyr Pro Ala Ser Gln Pro Pro Tyr Asn Pro Xaa
 115 120 125

Tyr Met Asp Ala Pro Lys Xaa Xaa Ser Glu His Ser Leu Ala Ser Leu
 130 135 140

45 Ala Ala Thr Trp Leu Cys Cys Val Cys Ala Xaa
 145 150 155

50 (2) INFORMATION FOR SEQ ID NO: 226:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 10 amino acids

(B) TYPE: amino acid

55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 226:

Met Gly Phe Gly Ala Thr Leu Ala Val Gly
 1 5 10

60

(2) INFORMATION FOR SEQ ID NO: 227:

5 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 20 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 227:

10 Met Ser Ile Phe Leu Val Met Ser Ile Ser Cys Ser Ser Thr Ser His
 1 5 10 15

15 Cys Tyr Ser Phe
 20

(2) INFORMATION FOR SEQ ID NO: 228:

20 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 94 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 228:

Met Ser Phe Ser Phe Ile Ile Phe Leu Leu Leu Val Cys Gln Glu Ile
 1 5 10 15

30 Thr Phe Cys Met Ser Tyr Gly Asp Ala Val Asn Cys Phe Ser Glu Cys
 20 25 30

Phe Ser Asn Leu Gln Thr Ile Tyr Ile Ser Cys Leu Gln His Ala Val
 35 40 45

35 Cys Lys His Ser Val Ile Trp Ser Ile Gln Leu Phe Val Arg Ala Leu
 50 55 60

40 Pro Ile Ser Lys Cys Ala Glu Leu Ser Ile Asp Gly Ile Phe Arg Ser
 65 70 75 80

Phe His Glu Asn Trp Lys Cys Ser Trp Val Ala Pro Thr Xaa
 85 90

(2) INFORMATION FOR SEQ ID NO: 229:

50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 94 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 229:

55 Met Ser Phe Ser Phe Ile Ile Phe Leu Leu Leu Val Cys Gln Glu Ile
 1 5 10 15

60 Thr Phe Cys Met Ser Tyr Gly Asp Ala Val Asn Cys Phe Ser Glu Cys
 20 25 30

336

Phe Ser Asn Leu Gln Thr Ile Tyr Ile Ser Cys Leu Gln His Ala Val
 35 40 45
 5 Cys Lys His Ser Val Ile Trp Ser Ile Gln Leu Phe Val Arg Ala Leu
 50 55 60
 Pro Ile Ser Lys Cys Ala Glu Leu Ser Ile Asp Gly Ile Phe Arg Ser
 65 70 75 80
 10 Phe His Glu Asn Trp Lys Cys Ser Trp Val Ala Pro Thr Xaa
 85 90

15 (2) INFORMATION FOR SEQ ID NO: 230:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 37 amino acids
 (B) TYPE: amino acid
 20 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 230:

Met Gly Trp Ser Ala Gly Leu Leu Phe Leu Leu Ile Leu Tyr Leu Pro
 1 5 10 15
 25 Val Pro Gly Trp Met Glu Arg Glu Asp Gly Gly Asp Gly Thr Ser Phe
 20 25 30
 Thr Ser Gly Ser Trp
 30 35

35 (2) INFORMATION FOR SEQ ID NO: 231:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 81 amino acids
 (B) TYPE: amino acid
 40 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 231:

Met Ala Thr Leu Trp Gly Gly Leu Leu Arg Leu Gly Ser Leu Leu Ser
 1 5 10 15
 45 Leu Ser Cys Leu Ala Leu Ser Val Leu Leu Leu Ala His Val Gln Thr
 20 25 30
 Pro Pro Arg Ile Ser Arg Met Ser Asp Val Ala Val Ser Ala Leu Pro
 35 40 45
 50 Ile Lys Lys Ile Leu Gly Ile Phe Ile Ile Ala Thr Tyr Leu Arg Lys
 50 55 60
 55 Ile Val Ile Ala Phe Met Leu Trp Ser Ser Val Leu Tyr Gly Gly Leu
 65 70 75 80
 Met

60

(2) INFORMATION FOR SEQ ID NO: 232:

(i) SEQUENCE CHARACTERISTICS:

5

(A) LENGTH: 301 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 232:

10 Met Asp Ala Arg Trp Trp Ala Val Val Val Leu Ala Ala Phe Pro Ser
 1 5 10 15
 Leu Gly Ala Gly Gly Glu Thr Pro Glu Ala Pro Pro Glu Ser Trp Thr
 20 25 30
 15 Gln Leu Trp Phe Phe Arg Phe Val Val Asn Ala Ala Gly Tyr Ala Xaa
 35 40 45
 Phe Met Val Pro Gly Tyr Leu Leu Val Gln Tyr Phe Arg Arg Lys Asn
 20 50 55 60
 Tyr Leu Glu Thr Gly Arg Gly Leu Cys Phe Pro Leu Val Lys Ala Cys
 65 70 75 80
 25 Val Phe Gly Asn Glu Pro Lys Ala Ser Asp Glu Val Pro Leu Ala Pro
 85 90 95
 Arg Thr Glu Ala Ala Glu Thr Thr Pro Met Trp Gln Ala Leu Lys Leu
 100 105 110
 30 Leu Phe Cys Ala Thr Gly Leu Gln Val Ser Tyr Leu Thr Trp Gly Val
 115 120 125
 Leu Gln Glu Arg Val Met Thr Arg Ser Tyr Gly Ala Thr Ala Thr Ser
 35 130 135 140
 Pro Gly Glu Arg Phe Thr Asp Ser Gln Phe Leu Val Leu Met Asn Arg
 145 150 155 160
 40 Val Leu Ala Leu Ile Val Ala Gly Leu Ser Cys Val Leu Cys Lys Gln
 165 170 175
 Pro Arg His Gly Ala Pro Met Tyr Arg Tyr Ser Phe Ala Ser Leu Ser
 180 185 190
 45 Asn Val Leu Ser Ser Trp Cys Gln Tyr Glu Ala Leu Lys Phe Val Ser
 195 200 205
 Phe Pro Thr Gln Val Leu Ala Lys Ala Ser Lys Val Ile Pro Val Met
 50 210 215 220
 Leu Met Gly Lys Leu Val Ser Arg Arg Xaa Asn Glu His Trp Glu Tyr
 225 230 235 240
 55 Leu Thr Ala Thr Leu Ile Ser Ile Gly Val Ser Met Phe Leu Leu Ser
 245 250 255
 Ser Gly Pro Glu Pro Arg Ser Ser Pro Ala Thr Thr Leu Ser Gly Leu
 260 265 270
 60

338

Ile Leu Leu Ala Gly Tyr Ile Ala Phe Asp Ser Phe Thr Ser Asn Trp
 275 280 285

5 Gln Asp Ala Cys Leu Pro Ile Arg Cys His Arg Cys Arg
 290 295 300

(2) INFORMATION FOR SEQ ID NO: 233:

10 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 313 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 233:

Met Ser Asp Leu Leu Leu Leu Gly Leu Ile Gly Gly Leu Thr Leu Leu
 1 5 10 15

20 Leu Leu Leu Thr Leu Leu Ala Phe Ala Gly Tyr Ser Gly Leu Leu Ala
 20 25 30

Gly Val Glu Val Ser Ala Gly Ser Pro Pro Ile Arg Asn Val Thr Val
 35 40 45

25 Ala Tyr Lys Phe His Met Gly Leu Tyr Gly Glu Thr Gly Arg Leu Phe
 50 55 60

30 Thr Glu Ser Cys Ser Ile Ser Pro Lys Leu Arg Ser Ile Ala Val Tyr
 65 70 75 80

Tyr Asp Asn Pro His Met Val Pro Pro Asp Lys Cys Arg Cys Ala Val
 85 90 95

35 Gly Ser Ile Leu Ser Glu Gly Glu Glu Ser Pro Ser Pro Glu Leu Ile
 100 105 110

Asp Leu Tyr Gln Lys Phe Gly Phe Lys Val Phe Ser Phe Pro Ala Pro
 115 120 125

40 Ser His Val Val Thr Ala Thr Phe Pro Tyr Thr Thr Ile Leu Ser Ile
 130 135 140

Trp Leu Ala Thr Arg Arg Val His Pro Ala Leu Asp Thr Tyr Ile Lys
 145 150 155 160

Glu Arg Lys Leu Cys Ala Tyr Pro Arg Leu Glu Ile Tyr Gln Glu Asp
 165 170 175

50 Gln Ile His Phe Met Cys Pro Leu Ala Xaa Gln Gly Asp Phe Tyr Val
 180 185 190

Pro Glu Met Lys Glu Thr Glu Trp Lys Trp Arg Gly Leu Val Glu Ala
 195 200 205

55 Ile Asp Thr Gln Val Asp Gly Thr Gly Ala Asp Thr Met Ser Asp Thr
 210 215 220

60 Ser Ser Val Ser Leu Glu Val Ser Pro Gly Ser Arg Glu Thr Ser Ala
 225 230 235 240

339

(2) INFORMATION FOR SEQ ID NO: 234:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 48 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 234:

Pro Gln Ser Leu Ile Leu His Leu Leu Leu Phe Phe Phe Leu Leu Phe
1 5 10 15

Leu Phe Phe Ile Phe Ile Phe Leu Phe Phe Leu Gln Cys Leu Thr Phe
20 25 30

Leu Phe Xaa Lys Pro Arg Gly Arg Tyr His Gly Leu Cys Phe Lys Phe
35 40 45

40 (2) INFORMATION FOR SEQ ID NO: 235:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 34 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 235:

50 Pro Ala Leu Arg Pro Ala Leu Leu Trp Ala Leu Leu Ala Leu Trp Leu
1 5 10 15

Cys Cys Ala Thr Pro Arg Met His Cys Ser Val Glu Met Ala Met Asn
20 25 30

55 Pro Val

60 (2) INFORMATION FOR SEQ ID NO: 236:

340

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 313 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 236:

Met Thr Arg Gly Gly Pro Gly Gly Arg Pro Gly Leu Pro Gln Pro Pro
 1 5 10 15
 Pro Leu Leu Leu Leu Leu Leu Xaa Leu Leu Leu Val Thr Ala Glu
 20 25 30
 Pro Pro Lys Pro Ala Gly Val Tyr Tyr Ala Thr Ala Tyr Trp Met Pro
 35 40 45
 Ala Glu Lys Thr Val Gln Val Lys Asn Val Met Asp Lys Asn Gly Asp
 50 55 60
 Ala Tyr Gly Phe Tyr Asn Asn Ser Val Lys Thr Thr Gly Trp Gly Ile
 65 70 75 80
 Leu Glu Ile Arg Ala Gly Tyr Gly Ser Gln Thr Leu Ser Asn Glu Ile
 85 90 95
 Ile Met Phe Val Ala Gly Phe Leu Glu Gly Tyr Leu Thr Ala Pro His
 100 105 110
 Met Asn Asp His Tyr Thr Asn Leu Tyr Pro Gln Leu Ile Thr Lys Pro
 115 120 125
 Ser Ile Met Asp Lys Val Gln Asp Phe Met Glu Lys Gln Asp Lys Trp
 130 135 140
 Thr Arg Lys Asn Ile Lys Glu Tyr Lys Thr Asp Ser Phe Trp Arg His
 145 150 155 160
 Thr Gly Tyr Val Met Ala Gln Ile Asp Gly Leu Tyr Val Gly Ala Lys
 165 170 175
 Lys Arg Ala Ile Leu Glu Gly Thr Lys Pro Met Thr Leu Phe Gln Ile
 180 185 190
 Gln Phe Leu Asn Ser Val Gly Asp Leu Leu Asp Leu Ile Pro Ser Leu
 195 200 205
 Ser Pro Thr Lys Asn Gly Ser Leu Lys Val Phe Lys Arg Trp Asp Met
 210 215 220
 Gly His Cys Ser Ala Leu Ile Lys Val Leu Pro Gly Phe Glu Asn Ile
 225 230 235 240
 Leu Phe Ala His Ser Ser Trp Tyr Thr Tyr Ala Ala Met Leu Arg Ile
 245 250 255
 Tyr Lys His Trp Asp Phe Asn Xaa Ile Asp Lys Asp Thr Ser Ser Ser
 260 265 270
 Arg Leu Ser Phe Ser Ser Tyr Pro Gly Phe Leu Glu Ser Leu Asp Asp
 275 280 285

341

Phe Tyr Ile Leu Ser Ser Gly Leu Ile Leu Leu Gln Thr Thr Asn Ser
 290 295 300

5 Val Phe Asn Lys Thr Leu Leu Lys Gln
 305 310

10 (2) INFORMATION FOR SEQ ID NO: 237:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 296 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 237:

Met Leu Gln Gly Pro Gly Ser Leu Leu Leu Leu Phe Leu Ala Ser His
 1 5 10 15

20 Cys Cys Leu Gly Ser Ala Arg Gly Leu Phe Leu Phe Gly Gln Pro Asp
 20 25 30

25 Phe Ser Tyr Lys Arg Xaa Asn Cys Lys Pro Ile Pro Val Asn Leu Gln
 35 40 45

Leu Cys His Gly Ile Glu Tyr Gln Asn Met Arg Leu Pro Asn Leu Leu
 50 55 60

30 Gly His Glu Thr Met Lys Glu Val Leu Glu Gln Ala Gly Ala Trp Ile
 65 70 75 80

Pro Leu Val Met Lys Gln Cys His Pro Asp Thr Lys Lys Phe Leu Cys
 85 90 95

35 Ser Leu Phe Ala Pro Val Cys Leu Asp Asp Leu Asp Glu Thr Ile Gln
 100 105 110

40 Pro Cys His Ser Leu Cys Val Gln Val Lys Asp Arg Cys Ala Pro Val
 115 120 125

Met Ser Ala Phe Gly Phe Pro Trp Pro Asp Met Leu Glu Cys Asp Arg
 130 135 140

45 Phe Pro Gln Asp Asn Asp Leu Cys Ile Pro Leu Ala Ser Ser Asp His
 145 150 155 160

Leu Leu Pro Ala Thr Glu Glu Ala Pro Lys Val Cys Glu Ala Cys Lys
 165 170 175

50 Asn Lys Asn Asp Asp Asp Asn Asp Ile Met Glu Thr Leu Cys Lys Asn
 180 185 190

55 Asp Phe Ala Leu Lys Ile Lys Val Lys Glu Ile Thr Tyr Ile Asn Arg
 195 200 205

Asp Thr Lys Ile Ile Leu Glu Thr Lys Ser Lys Thr Ile Tyr Lys Leu
 210 215 220

60 Asn Gly Val Ser Glu Arg Asp Leu Lys Lys Ser Val Leu Trp Leu Lys

342

225 230 235 240
 Asp Ser Leu Gln Cys Thr Cys Glu Glu Met Asn Asp Ile Asn Ala Pro
 245 250 255
 5 Tyr Leu Val Met Gly Gln Lys Gln Gly Gly Glu Leu Val Ile Thr Ser
 260 265 270
 10 Val Lys Arg Trp Gln Lys Gly Gln Arg Glu Phe Lys Arg Ile Ser Arg
 275 280 285
 Ser Ile Arg Lys Leu Gln Cys Xaa
 290 295

15

(2) INFORMATION FOR SEQ ID NO: 238:

(i) SEQUENCE CHARACTERISTICS:
 20 (A) LENGTH: 92 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 238:

25 Met Ala Ser Leu Gly His Ile Leu Val Phe Cys Val Gly Leu Leu Thr
 1 5 10 15
 Met Ala Lys Ala Glu Ser Pro Lys Glu His Asp Pro Phe Thr Tyr Asp
 20 25 30
 30 Tyr Gln Ser Leu Gln Ile Gly Gly Leu Val Ile Ala Gly Ile Leu Phe
 35 40 45
 Ile Leu Gly Ile Leu Ile Val Leu Ser Arg Arg Cys Arg Cys Lys Phe
 35 50 55 60
 Asn Gln Gln Gln Arg Thr Gly Glu Pro Asp Glu Glu Gly Thr Phe
 65 70 75 80
 40 Arg Ser Ser Ile Arg Arg Leu Ser Xaa Arg Xaa Arg
 85 90

45 (2) INFORMATION FOR SEQ ID NO: 239:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 71 amino acids
 (B) TYPE: amino acid
 50 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 239:

Met Pro Gly Thr Phe Leu Arg Pro Phe Val Phe Leu Phe Leu Phe Ile
 1 5 10 15
 55 Cys Cys Cys Leu His Ser Gly Gly Leu Gly Gly Val Pro Leu Pro Pro
 20 25 30
 Phe Pro Pro Gln Ala Gln Arg Gly Glu Gly Pro Gly Lys Trp Met Ser
 60 35 40 45

343

Pro Pro Leu Pro Pro His Pro Val Val Ala Pro Pro Thr Pro Ser Pro
 50 55 60

5 Ser Arg Gly Cys Val Leu Leu
 65 70

10 (2) INFORMATION FOR SEQ ID NO: 240:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 71 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 240:

Met Pro Gly Thr Phe Leu Arg Pro Phe Val Phe Leu Phe Leu Phe Ile
 1 5 10 15

20 Cys Cys Cys Leu His Ser Gly Gly Leu Gly Gly Val Pro Leu Pro Pro
 20 25 30

25 Phe Pro Pro Gln Ala Gln Arg Gly Glu Gly Pro Gly Lys Trp Met Ser
 35 40 45

Pro Pro Leu Pro Pro His Pro Val Val Ala Pro Pro Thr Pro Ser Pro
 50 55 60

30 Ser Arg Gly Cys Val Leu Leu
 65 70

35 (2) INFORMATION FOR SEQ ID NO: 241:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 28 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 241:

Met Phe Tyr Val Leu Ser Val Ser Xaa Leu Xaa Leu Phe Leu Ala Cys
 1 5 10 15

45 Gly Leu Cys Leu Xaa Leu Leu Thr Gly Lys Leu Leu
 20 25

50

(2) INFORMATION FOR SEQ ID NO: 242:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 58 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 242:

Met Lys Leu Phe Asp Ala Ser Pro Thr Phe Phe Ala Phe Leu Leu Gly
 1 5 10 15

60

344

His Ile Leu Ala Met Glu Val Leu Ala Trp Leu Leu Ile Tyr Leu Leu
 20 25 30

5 Gly Pro Gly Trp Val Pro Ser Ala Leu Xaa Arg Leu His Pro Gly His
 35 40 45

Leu Ser Gly Ser Val Leu Val Ser Ala Ala
 50 55

10

(2) INFORMATION FOR SEQ ID NO: 243:

15 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 123 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 243:

20 Met Ile Leu Gly Gly Ile Val Val Val Leu Val Phe Thr Gly Phe Val
 1 5 10 15

25 Trp Ala Ala His Asn Lys Asp Val Leu Arg Arg Met Lys Lys Arg Tyr
 20 25 30

Pro Thr Thr Phe Val Met Val Val Met Leu Ala Ser Tyr Phe Leu Ile
 35 40 45

30 Ser Met Phe Gly Gly Val Met Val Phe Val Phe Gly Ile Thr Phe Pro
 50 55 60

Leu Leu Leu Met Phe Ile His Ala Ser Leu Arg Leu Arg Asn Leu Lys
 65 70 75 80

35 Asn Lys Leu Glu Asn Lys Met Glu Gly Ile Gly Leu Lys Arg Thr Pro
 85 90 95

Met Gly Ile Val Leu Asp Ala Leu Glu Gln Gln Glu Glu Gly Ile Asn
 100 105 110

40 Arg Leu Thr Asp Tyr Ile Ser Lys Val Lys Glu
 115 120

45

(2) INFORMATION FOR SEQ ID NO: 244:

50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 73 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 244:

55 Ala Leu Val Ser Gly Gln Leu Cys Met Glu Ile Ala Arg Gly Asn Ile
 1 5 10 15

Phe Phe Leu Asn Xaa Leu Val Thr Thr Phe Tyr Cys Ser Cys Leu Leu
 20 25 30

60

345

Leu Ser Val Xaa Tyr Leu His Xaa Gly Phe Phe Tyr Ser Ser Leu Cys
 35 40 45

5 Lys Cys Cys Phe Val Leu Val Val Leu Ser Arg Ile Gly Ser Val Asn
 50 55 60

Glu Thr Trp Ser Cys Asn Phe Ser Ile
 65 70

10

(2) INFORMATION FOR SEQ ID NO: 245:

15 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 49 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 245:

20 Thr Pro Ala Thr Thr Ser Ser Ser Ser Ser Pro Leu Phe Leu Ser Ser
 1 5 10 15

Pro Asp Trp Ser Ser Cys Pro Ser Gly Ser Cys Ile Ala Pro Trp Cys
 20 25 30

25 Thr His Trp Ser Ser Ile Leu Pro Ser Leu Xaa Ile Thr Ser Ser Ile
 35 40 45

Pro
 30

(2) INFORMATION FOR SEQ ID NO: 246:

35 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 339 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 246:

Met Ala Arg Val Pro Pro Leu Ser Ser Ser Trp Thr Ser Ser Arg Tyr
 1 5 10 15

45 Arg Arg Trp Leu Cys Cys Pro Val Trp Trp Thr Thr Phe Trp Ala Thr
 20 25 30

Ala Trp Ser Leu Thr Lys His Leu Tyr Lys Asp Val Thr Asp Ala Ile
 35 40 45

50 Arg Asp Val His Val Lys Gly Leu Met Tyr Ser Trp Ile Glu Gln Asp
 50 55 60

55 Met Glu Lys Tyr Ile Leu Arg Gly Asp Glu Ile Phe Ala Val Leu Ser
 65 70 80

Arg Leu Val Ala His Gly Lys Gln Leu Phe Thr Thr Thr Asn Ser Pro
 85 90 95

60 Phe Ser Phe Val Asp Lys Gly Met Arg His Met Val Gly Pro Asp Trp

346

	100	105	110
5	Arg His Ser Ser Met Trp Ser Leu Ser Arg Gln Thr Ser Pro Ala Ser 115 120 125		
	Ser Leu Thr Gly Ala Thr Phe Arg Lys Leu Asp Glu Lys Gly Ser Leu 130 135 140		
10	Gln Trp Asp Arg Ile Thr Arg Leu Glu Lys Gly Lys Ile Tyr Arg Gln 145 150 155 160		
	Gly Asn Leu Phe Asp Phe Leu Arg Leu Thr Glu Trp Arg Gly Pro Arg 165 170 175		
15	Val Leu Tyr Phe Gly Asp His Leu Tyr Ser Asp Leu Ala Asp Leu Met 180 185 190		
	Leu Arg His Gly Trp Arg Thr Gly Ala Ile Ile Pro Glu Leu Glu Arg 195 200 205		
20	Glu Ile Arg Ile Ile Asn Thr Glu Gln Tyr Met His Ser Leu Thr Trp 210 215 220		
	Gln Gln Ala Leu Thr Gly Leu Leu Glu Arg Met Gln Thr Tyr Gln Asp 225 230 235 240		
25	Ala Glu Ser Arg Gln Val Leu Ala Ala Trp Met Lys Glu Arg Gln Glu 245 250 255		
	Leu Arg Cys Ile Thr Lys Ala Leu Phe Asn Ala Gln Phe Gly Ser Ile 260 265 270		
30	Phe Arg Thr Phe His Asn Pro Thr Tyr Phe Ser Arg Arg Leu Val Arg 275 280 285		
35	Phe Ser Asp Leu Tyr Met Ala Ser Leu Ser Cys Leu Leu Asn Tyr Arg 290 295 300		
	Val Asp Phe Thr Phe Tyr Pro Arg Arg Thr Pro Leu Gln His Glu Ala 305 310 315 320		
40	Pro Leu Trp Met Asp Gln Leu Leu His Arg Leu His Glu Asp Pro Leu 325 330 335		
45	Pro Trp Xaa		

50 (2) INFORMATION FOR SEQ ID NO: 247:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 18 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID: 1

Met Ala Leu Leu Ser Cys Val Val Asp Tyr Thr Leu Gly His Ser Leu
1 5 10 15

60

Xaa Val

5

(2) INFORMATION FOR SEQ ID NO: 248:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 339 amino acids

10

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 248:

15

Met Asn Trp Glu Leu Leu Leu Trp Leu Leu Val Leu Cys Ala Leu Leu
1 5 10 15Leu Leu Leu Val Gln Leu Leu Arg Phe Leu Arg Ala Asp Gly Asp Leu
20 25 30

20

Thr Leu Leu Trp Ala Glu Trp Gln Gly Arg Arg Pro Glu Trp Glu Leu
35 40 45Thr Asp Met Val Val Trp Val Thr Gly Ala Ser Ser Gly Ile Gly Glu
50 55 60

25

Glu Leu Ala Tyr Gln Leu Ser Lys Leu Gly Val Ser Leu Val Leu Ser
65 70 75 80Ala Arg Arg Val His Glu Leu Glu Arg Val Lys Arg Arg Cys Leu Glu
85 90 95

30

Asn Gly Asn Leu Lys Glu Lys Asp Ile Leu Val Leu Pro Leu Asp Leu
100 105 110

35

Thr Asp Thr Gly Ser His Glu Ala Ala Thr Lys Ala Val Leu Gln Glu
115 120 125Phe Gly Arg Ile Asp Ile Leu Val Asn Asn Gly Gly Met Ser Gln Arg
130 135 140

40

Ser Leu Cys Met Asp Thr Ser Leu Asp Val Tyr Arg Lys Leu Ile Glu
145 150 155 160Leu Asn Tyr Leu Gly Thr Val Ser Leu Thr Lys Cys Val Leu Pro His
165 170 175

45

Met Ile Glu Arg Lys Gln Gly Lys Ile Val Thr Val Asn Ser Ile Leu
180 185 190

50

Gly Ile Ile Ser Val Pro Leu Ser Ile Gly Tyr Cys Ala Ser Lys His
195 200 205Ala Leu Arg Gly Phe Phe Asn Gly Leu Arg Thr Glu Leu Ala Thr Tyr
210 215 220

55

Pro Gly Ile Ile Val Ser Asn Ile Cys Pro Gly Pro Val Gln Ser Asn
225 230 235 240Ile Val Glu Asn Ser Leu Ala Gly Glu Val Thr Lys Thr Ile Gly Asn
245 250 255

60

348

Asn Gly Asp Gln Ser His Lys Met Thr Thr Ser Arg Cys Val Arg Leu
 260 265 270

5 Met Leu Ile Ser Met Ala Asn Asp Leu Lys Glu Val Trp Ile Ser Glu
 275 280 285

Gln Pro Phe Leu Leu Val Thr Tyr Leu Trp Gln Tyr Met Pro Thr Trp
 290 295 300

10 Ala Trp Trp Ile Thr Asn Lys Met Gly Lys Lys Arg Ile Glu Asn Phe
 305 310 315 320

Lys Ser Gly Val Asp Ala Asp Ser Ser Tyr Phe Lys Ile Phe Lys Thr
 15 325 330 335

Lys His Asp

20

(2) INFORMATION FOR SEQ ID NO: 249:

25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 96 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 249:

30 Met Gly Ala Arg Pro Gly Gly His Pro Gln Lys Trp Ser Phe Leu Trp
 1 5 10 15

Ser Leu Ala Leu Trp Leu Pro Leu Ala Leu Ser Val Ser Leu Phe Leu
 20 25 30

35 Gly Leu Ser Leu Ser Pro Pro Gln Pro Gly Leu Ser Leu Trp Cys Thr
 35 40 45

40 Leu Ser Tyr Cys Cys Glu Gln Trp Lys Phe Lys Gly Thr Pro Ser Pro
 50 55 60

Ala Leu Leu Asn Leu Gly Thr Gln Pro Lys Lys Asp Lys Lys Leu Glu
 65 70 75 80

45 Asp Ser Ile Ala Thr Gln Leu Arg Xaa Leu Pro Glu Lys Asn Ser Asn
 85 90 95

50

(2) INFORMATION FOR SEQ ID NO: 250:

55 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 79 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 250:

60

349

Met Ala Leu Thr Phe Leu Leu Val Leu Leu Thr Leu Ala Thr Leu Cys
 1 5 10 15

5 Thr Arg Leu His Arg Asn Phe Arg Arg Gly Glu Ser Ile Tyr Trp Gly
 20 25 30

Pro Thr Ala Asp Ser Gln Asp Thr Val Ala Ala Val Leu Lys Arg Arg
 35 40 45

10 Leu Leu Gln Pro Ser Arg Arg Val Lys Arg Ser Arg Arg Arg Pro Xaa
 50 55 60

Xaa Pro Pro Thr Pro Asp Ser Gly Pro Glu Gly Glu Ser Ser Glu
 65 70 75

15

(2) INFORMATION FOR SEQ ID NO: 251:

20 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 354 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 251:

25 Met Gly Pro Ser Thr Pro Leu Leu Ile Leu Phe Leu Leu Ser Trp Ser
 1 5 10 15

30 Gly Pro Leu Gln Gly Gln Gln His His Leu Val Glu Tyr Met Glu Arg
 20 25 30

Arg Leu Ala Ala Leu Glu Glu Arg Leu Ala Gln Cys Gln Asp Gln Ser
 35 40 45

35 Ser Arg His Ala Ala Glu Leu Arg Asp Phe Lys Asn Lys Met Leu Pro
 50 55 60

Leu Leu Glu Val Ala Glu Lys Glu Arg Glu Ala Leu Arg Thr Glu Ala
 65 70 75 80

40 Asp Thr Ile Ser Gly Arg Val Asp Arg Leu Glu Arg Glu Val Asp Tyr
 85 90 95

Leu Glu Thr Gln Asn Pro Ala Leu Pro Cys Val Glu Phe Asp Glu Lys
 45 100 105 110

Val Thr Gly Gly Pro Gly Thr Lys Gly Lys Gly Arg Arg Asn Glu Lys
 115 120 125

50 Tyr Asp Met Val Thr Asp Cys Gly Tyr Thr Ile Ser Gln Val Arg Ser
 130 135 140

Met Lys Ile Leu Lys Arg Phe Gly Gly Pro Ala Gly Leu Trp Thr Lys
 145 150 155 160

55 Asp Pro Leu Gly Gln Thr Glu Lys Ile Tyr Val Leu Asp Gly Thr Gln
 165 170 175

Asn Asp Thr Ala Phe Val Phe Pro Arg Leu Arg Asp Phe Thr Leu Ala
 60 180 185 190

350

Met Ala Ala Arg Lys Ala Ser Arg Val Arg Val Pro Phe Pro Trp Val
 195 200 205

5 Gly Thr Gly Gln Leu Val Tyr Gly Gly Phe Leu Tyr Phe Ala Arg Arg
 210 215 220

Pro Pro Gly Arg Pro Gly Gly Gly Gly Glu Met Glu Asn Thr Leu Gln
 225 230 235 240

10 Leu Ile Lys Phe His Leu Ala Asn Arg Thr Val Val Asp Ser Ser Val
 245 250 255

15 Phe Pro Ala Glu Gly Leu Ile Pro Pro Tyr Gly Leu Thr Ala Asp Thr
 260 265 270

Tyr Ile Asp Leu Ala Ala Asp Glu Glu Gly Leu Trp Ala Val Tyr Ala
 275 280 285

20 Thr Arg Glu Asp Asp Arg His Leu Cys Leu Ala Lys Leu Asp Pro Gln
 290 295 300

Thr Leu Asp Thr Glu Gln Gln Trp Asp Thr Pro Cys Pro Arg Glu Asn
 305 310 315 320

25 Ala Glu Ala Ala Phe Xaa Ile Cys Gly Thr Leu Tyr Val Val Tyr Asn
 325 330 335

30 Thr Arg Pro Ala Ser Arg Ala Arg Ile Gln Cys Ser Phe Asp Ala Ser
 340 345 350

Gly Pro

35

(2) INFORMATION FOR SEQ ID NO: 252:

- (i) SEQUENCE CHARACTERISTICS:
- 40 (A) LENGTH: 109 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 252:

45 Met Leu Cys Ile Asn Gly Thr Thr Pro Arg Pro Leu Pro Val Pro Ser
 1 5 10 15

Pro Phe Gly Cys Met Ile Phe Phe Phe Phe Lys Asn Pro Trp Lys Gln
 20 25 30

50 Arg Leu Leu Gln Gly Trp Leu Gly Ala Arg Pro Ile His Leu Leu Gly
 35 40 45

Tyr Leu Pro Leu Ser Leu Leu Trp Cys Pro Phe Pro Leu Pro Cys Ala
 50 55 60

Arg Cys Ser Val Val Tyr Ile Ser Ser Pro Arg His Gly Ala His Ala
 65 70 75 80

60 Pro Arg Asp Met Ile Leu Ser Leu Val Leu Ala His Gly Ala Leu Tyr

351

85 90 95

Lys Glu Leu Gly Gly Arg Gly Arg Lys Trp Glu Pro Ser
100 105

5

(2) INFORMATION FOR SEQ ID NO: 253:

10 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 45 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 253:

15 Met Phe Tyr Phe Leu Pro Leu Ile Phe Pro Ala Phe Pro Pro Trp Ala
1 5 10 15

20 Phe Arg Leu Ser Thr Leu Phe Thr Ile Ile Ser Trp Ser Glu Asp Ser
20 25 30

Asn Asn Ser Gln Val Tyr Met Asn Cys Val Cys Ser Phe
35 40 45

25

(2) INFORMATION FOR SEQ ID NO: 254:

30 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 315 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 254:

35 Met Ala Gly Gly Arg Cys Gly Pro Xaa Leu Thr Ala Leu Leu Ala Ala
1 5 10 15

Trp Ile Ala Ala Val Ala Ala Thr Ala Gly Pro Glu Glu Ala Ala Leu
20 25 30

40 Pro Pro Glu Gln Ser Arg Val Gln Pro Met Thr Ala Ser Asn Trp Thr
35 40 45

45 Leu Val Met Glu Gly Glu Trp Met Leu Lys Phe Tyr Ala Pro Trp Cys
50 55 60

Pro Ser Cys Gln Gln Thr Asp Ser Glu Trp Glu Ala Phe Ala Lys Asn
65 70 75 80

50 Gly Glu Ile Leu Gln Ile Ser Val Gly Lys Val Asp Val Ile Gln Glu
85 90 95

Pro Gly Leu Ser Gly Arg Phe Phe Val Thr Thr Leu Pro Ala Phe Phe
100 105 110

55 His Ala Lys Asp Gly Ile Phe Arg Arg Tyr Arg Gly Pro Gly Ile Phe
115 120 125

60 Glu Asp Leu Gln Asn Tyr Ile Leu Glu Lys Lys Trp Gln Ser Val Glu
130 135 140

352

Pro Leu Thr Gly Trp Lys Ser Pro Ala Ser Leu Thr Met Ser Gly Met
 145 150 155 160

5 Ala Gly Leu Phe Ser Ile Ser Gly Lys Ile Trp His Leu His Asn Tyr
 165 170 175

Phe Thr Val Thr Leu Gly Ile Pro Ala Trp Cys Ser Tyr Val Phe Phe
 180 185 190

10 Val Ile Ala Thr Leu Val Phe Gly Leu Phe Met Gly Leu Val Leu Val
 195 200 205

Val Ile Ser Glu Cys Phe Tyr Val Pro Leu Pro Arg His Leu Ser Glu
 15 210 215 220

Arg Ser Glu Gln Asn Arg Arg Ser Glu Glu Ala His Arg Ala Glu Gln
 225 230 235 240

20 Leu Gln Asp Ala Glu Glu Glu Lys Asp Asp Ser Asn Glu Glu Glu Asn
 245 250 255

Lys Asp Ser Leu Val Asp Asp Glu Glu Glu Lys Glu Asp Leu Gly Asp
 260 265 270

25 Glu Asp Glu Ala Glu Glu Glu Glu Glu Glu Asp Asn Leu Ala Ala Gly
 275 280 285

Val Asp Glu Glu Arg Ser Glu Ala Asn Asp Gln Gly Pro Pro Gly Glu
 30 290 295 300

Asp Gly Val Thr Arg Glu Xaa Ser Arg Ala Xaa
 305 310 315

35

(2) INFORMATION FOR SEQ ID NO: 255:

40 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 53 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 255:

45 Met Leu Lys Ala Leu Phe Arg Thr Leu Gln Ala Met Leu Leu Gly Val
 1 5 10 15

Trp Ile Leu Leu Leu Ala Ser Leu Ala Pro Leu Trp Leu Tyr Cys
 20 25 30

50 Trp Arg Met Phe Pro Thr Lys Gly Lys Arg Asp Gln Lys Glu Met Leu
 35 40 45

Glu Val Ser Gly Ile
 55 50

60 (2) INFORMATION FOR SEQ ID NO: 256:

353

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 93 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 256:

Met Ile His Leu Gly His Ile Leu Phe Leu Leu Leu Leu Pro Val Ala
 1 5 10 15

10 Ala Ala Gln Thr Thr Pro Gly Glu Arg Ser Ser Leu Pro Ala Phe Tyr
 20 25 30

Pro Gly Thr Ser Gly Ser Cys Ser Gly Cys Gly Ser Leu Ser Leu Pro
 35 40 45

15 Leu Leu Ala Gly Leu Val Ala Ala Asp Ala Val Ala Ser Leu Leu Ile
 50 55 60

Val Gly Ala Val Phe Leu Cys Ala Arg Pro Arg Arg Ser Pro Ala Gln
 20 65 70 75 80

Asp Gly Lys Val Tyr Ile Asn Met Pro Gly Arg Gly Xaa
 85 90

25

(2) INFORMATION FOR SEQ ID NO: 257:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 12 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 257:

35 Pro Gly His Leu Leu Pro His Lys Trp Glu Asn Cys
 1 5 10

40 (2) INFORMATION FOR SEQ ID NO: 258:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1852 base pairs

(B) TYPE: nucleic acid

45 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 258:

50 TGGCATCTGT GAGCAGCTGC CAGGCTCCGG CCAGGATCCC TTCCTTCTCC TCATTGGCTG 60

ATGGATCCCA AGGGGCTCCT CTCCTTGACC TTCGTGCTGT TTCTCTCCCT GGCTTTTGGG 120

GCAAGCTACG GAACAGGTGG GCGCATGATG AACTGCCGAA AGATTCTCCG GCAGTTGGGA 180

55 AGCAAAGTGC TGCTGCCCT GACATATGAA AGGATAAATA AGAGCATGAA CAAAAGCATC 240

CACATTGTCTG TCACAATGGC AAAATCACTG GAGAACAGTG TCGAGAACAA AATAGTGTCT 300

60 CTTGATCCAT CCGAAGCAGG CCTCCACGT TATCTAGGAG ATCCCTACAA GTTTTATCTG 360

	GAGAATCTCA CCCTGGGGAT ACGGGAAAGC AGGAAGGAGG ATGAGGGATG GTACCTTATG	420
5	ACCCTGGAGA AAAATGTTTC AGTTCAGCGC TTTTGCCTGC AGTTGAGGCT TTATGAGCAG	480
	GTCTCCACTC CAGAAATTAA AGTTTTAAAC AAGACCCAGG AGAACGGGAC CTGCACCTTG	540
	ATACTGGGCT GCACAGTGGA GAAGGGGGAC CATGTGGCTT ACAGCTGGAG TGAAAAGGCG	600
10	GGCACCCACC CACTGAACCC AGCCAACAGC TCCCACCTCC TGTCCCTCAC CCTCGGCCCC	660
	CAGCATGCTG ACAATATCTA CATCTGCACC GTGAGCAACC CTATCAGCAA CAATTCCCAG	720
15	ACCTTCAGCC CGTGGCCCGG ATGCAGGACA GACCCCTCAG AAACAAAACC ATGGGCAGTG	780
	TATGCTGGGC TGTTAGGGGG TGTCATCATG ATTCTCATCA TGGTGGTAAT ACTACAGTTG	840
	AGAAGAAGAG GTAAAACGAA CCATTACCAG ACAACAGTGG AAAAAAAG CCTTACGATC	900
20	TATGCCCAAG TCCAGAAACC AGGTGACACT CATCATCAGA CTTCGGACTT ATTCTAATCC	960
	AGGATGACCT TATTTTGAAA TCCTTATCTT GACATCTGTG AAGACCTTTA TTCAAATAAA	1020
25	GTCACATTTT GACATTCTGC GAGGGGCTGG AGCCGGGCGG GGGCGATGTG GAGCGCGGGC	1080
	CGCGGCGGGG CTGCCTGGCC GGTGCTGTIG GGGCTGCTGC TGGCGCTGTT AGTGCCGGGC	1140
	GGTGGTGCCG CCAAGACCGG TGCGGAGCTC GTGACTGCGG GTCGGTGCTG AAGCTGCTCA	1200
30	ATACGCACCA CCGGTGCGGC TGCACTCGCA CGACATCAAA TACGGATCCG GCAGCGGCCA	1260
	GCAATCGGTG ACCGGCGTAG AGGTGCGAGC GACGAATAGC TACTGGCGGA TCCGCGGCGG	1320
35	CTCGGAGGGG GGTGCCCCGG CGGGTCCCCG GTGCGCTGCG GGCAGGCGGT GAGGTCACAC	1380
	ATGTGCTTAC GGGCAAGAAC CTGCACACGC ACCACTTCCC GTCGCCGCTG TCCAACAACC	1440
	AGGAAGTGAG TGCCAAAGGG GAAGACGGCG AGGGCGACGA CCTGGACCTA TGGACAGTGC	1500
40	GCTGCTCTGC TCTGGACAGC ACTGGGAGCG TGAGGCTGCT GTGGCGCCTT CCAGCATGTG	1560
	GCACCTCTGT GGTTCCTGTC AGTCACGGTA GCAGTATGGA AGCCCCATCC GTGGGCAGCA	1620
45	TGAGGTCCAC GCATGCCCAG TGCCAACACG CACAATACGT GGAAGGCCAT GGAAGGCATC	1680
	TTCATCAAGC CTAGTGTTGA GCCCTCTGCA GGTACGATG AACTCTGAGT GTGTGGATGG	1740
	ATGGGTGGAT GGAGGTGGC AGGTGGGGCG TCTGCAGGGC CACTCTTGGC AGAGACTTTG	1800
50	GGTTTGTAGG GGTCTCAAG TGCCTTTGTG ATTAAAGAAT GTTGGTCTAT GA	1852

55 (2) INFORMATION FOR SEQ ID NO: 259:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 371 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

60

355

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 259:

Met Glu Leu Glu Leu Asp Ala Gly Asp Gln Asp Leu Leu Ala Phe Leu
 1 5 10 15
 5 Leu Glu Glu Ser Gly Asp Leu Gly Thr Ala Pro Asp Glu Ala Val Arg
 20 25 30
 10 Ala Pro Leu Asp Trp Ala Leu Pro Leu Ser Glu Val Pro Ser Asp Trp
 35 40 45
 Glu Val Asp Asp Leu Leu Cys Ser Leu Leu Ser Pro Pro Ala Ser Leu
 50 55 60
 15 Asn Ile Leu Ser Ser Ser Asn Pro Cys Leu Val His His Asp His Thr
 65 70 75 80
 Tyr Ser Leu Pro Arg Glu Thr Val Ser Met Asp Leu Glu Ser Glu Ser
 85 90 95
 20 Cys Arg Lys Glu Gly Thr Gln Met Thr Pro Gln His Met Glu Glu Leu
 100 105 110
 25 Ala Glu Gln Glu Ile Ala Arg Leu Val Leu Thr Asp Glu Glu Lys Ser
 115 120 125
 Leu Leu Glu Lys Glu Gly Leu Ile Leu Pro Glu Thr Leu Pro Leu Thr
 130 135 140
 30 Lys Thr Glu Glu Gln Ile Leu Lys Arg Val Arg Arg Lys Ile Arg Asn
 145 150 155 160
 Lys Arg Ser Ala Gln Glu Ser Arg Arg Lys Lys Lys Val Tyr Val Gly
 165 170 175
 35 Gly Leu Glu Ser Arg Val Leu Lys Tyr Thr Ala Gln Asn Met Glu Leu
 180 185 190
 40 Gln Asn Lys Val Gln Leu Leu Glu Glu Gln Asn Leu Ser Leu Leu Asp
 195 200 205
 Gln Leu Arg Lys Leu Gln Ala Met Val Ile Glu Ile Ser Asn Lys Thr
 210 215 220
 45 Ser Ser Ser Ser Thr Cys Ile Leu Val Leu Leu Val Ser Phe Cys Leu
 225 230 235 240
 Leu Leu Val Pro Ala Met Tyr Ser Ser Asp Thr Arg Gly Ser Leu Pro
 245 250 255
 50 Ala Glu His Gly Val Leu Ser Arg Gln Leu Asn Ala Leu Pro Ser Glu
 260 265 270
 55 Asp Pro Tyr Gln Leu Glu Leu Pro Ala Leu Glu Ser Glu Val Pro Lys
 275 280 285
 Asp Ser Thr His Gln Trp Leu Asp Gly Ser Arg Lys Val Leu Gln Ala
 290 295 300
 60 Pro Gly Asn Thr Ser Cys Leu Leu His Tyr Ser Pro Gln Ala Pro Ser

356

305 310 315 320
 Ala Glu Pro Pro Leu Glu Trp Pro Phe Pro Asp Leu Ser Ser Glu Pro
 325 330 335
 5 Leu Cys Arg Gly Pro Ile Leu Pro Leu Gln Ala Asn Leu Thr Arg Lys
 340 345 350
 10 Gly Gly Trp Leu Pro Thr Gly Ser Pro Ser Val Ile Leu Gln Asp Arg
 355 360 365
 Tyr Ser Gly
 370

15

(2) INFORMATION FOR SEQ ID NO: 260:

- 20 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 12 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 260:

25 Cys Arg Cys Ala Ser Gly Phe Thr Gly Glu Asp Cys
 1 5 10

30 (2) INFORMATION FOR SEQ ID NO: 261:

- 35 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 12 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 261:

40 Cys Thr Cys Gln Val Gly Phe Thr Gly Lys Glu Cys
 1 5 10

(2) INFORMATION FOR SEQ ID NO: 262:

- 45 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 12 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 262:

50 Cys Leu Asn Leu Pro Gly Ser Tyr Gln Cys Gln Cys
 1 5 10

55

(2) INFORMATION FOR SEQ ID NO: 263:

- 60 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 12 amino acids
 (B) TYPE: amino acid

357

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 263:

5 Cys Lys Cys Leu Thr Gly Phe Thr Gly Gln Lys Cys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 264:

10

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 12 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 264:

Cys Gln Cys Leu Gln Gly Phe Thr Gly Gln Tyr Cys
1 5 10

20

(2) INFORMATION FOR SEQ ID NO: 265:

(i) SEQUENCE CHARACTERISTICS:

25

(A) LENGTH: 127 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 265:

30 Gly Leu Ala Cys Trp Leu Ala Gly Val Ile Phe Ile Asp Arg Lys Arg
1 5 10 15

Thr Gly Asp Ala Ile Ser Val Met Ser Glu Val Ala Gln Thr Leu Leu
20 25 30

35

Thr Gln Asp Val Xaa Val Trp Val Phe Pro Glu Gly Thr Arg Asn His
35 40 45

40 Asn Gly Ser Met Leu Pro Phe Lys Arg Gly Ala Phe His Leu Ala Val
50 55 60

Gln Ala Gln Val Pro Ile Val Pro Ile Val Met Ser Ser Tyr Gln Asp
65 70 75 80

45 Phe Tyr Cys Lys Lys Glu Arg Arg Phe Thr Ser Gly Gln Cys Gln Val
85 90 95

Arg Val Leu Pro Pro Val Pro Thr Glu Gly Leu Thr Pro Asp Asp Val
100 105 110

50

Pro Ala Leu Ala Asp Arg Val Arg His Ser Met Leu His Cys Phe
115 120 125

55

(2) INFORMATION FOR SEQ ID NO: 266:

(i) SEQUENCE CHARACTERISTICS:

60

(A) LENGTH: 98 amino acids

(B) TYPE: amino acid

358

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 266:

5 Pro Ser Ala Lys Tyr Phe Phe Lys Met Ala Phe Tyr Asn Gly Trp Ile
 1 5 10 15
 Leu Phe Leu Ala Val Leu Ala Ile Pro Val Cys Ala Val Arg Gly Arg
 20 25 30
 10 Asn Val Glu Asn Met Lys Ile Leu Arg Leu Met Leu Leu His Ile Lys
 35 40 45
 Tyr Leu Tyr Gly Ile Arg Val Glu Val Arg Gly Ala His His Phe Pro
 50 55 60
 15 Pro Ser Gln Pro Tyr Val Val Val Ser Asn His Gln Ser Ser Leu Asp
 65 70 75 80
 Leu Leu Gly Met Met Glu Val Leu Pro Gly Arg Cys Val Pro Ile Ala
 20 85 90 95
 Lys Arg

25

(2) INFORMATION FOR SEQ ID NO: 267:

30 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 9 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 267:

35 Thr Val Phe Arg Glu Ile Ser Thr Asp
 1 5

40 (2) INFORMATION FOR SEQ ID NO: 268:

45 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 11 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 268:

50 Leu Trp Ala Gly Ser Ala Gly Trp Pro Ala Gly
 1 5 10

(2) INFORMATION FOR SEQ ID NO: 269:

55 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 29 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 269:

60

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Ser Ile Leu Gly Ile Ile Ser Val Pro Leu Ser Ile Gly Tyr Cys Ala
 1 5 10 15

5 Ser Lys His Ala Leu Arg Gly Phe Phe Asn Gly Leu Arg
 20 25

10 (2) INFORMATION FOR SEQ ID NO: 270:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 8 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 270:

Met Ala Tyr His Gly Leu Thr Val
 1 5

20

(2) INFORMATION FOR SEQ ID NO: 271:

25 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 6 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 271:

30 Ile Ser Ala Ala Arg Val
 1 5

35 (2) INFORMATION FOR SEQ ID NO: 272:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 11 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 272:

Pro Asp Val Ser Glu Phe Met Thr Arg Leu Phe
 1 5 10

45

(2) INFORMATION FOR SEQ ID NO: 273:

50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 17 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 273:

55 Phe Asp Pro Val Arg Val Asp Ile Thr Ser Lys Gly Lys Met Arg Ala
 1 5 10 15

Arg


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Applicant's or agent's file reference number	P001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97901
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer


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Applicant's or agent's file reference number	PS001PCT	International application.	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64 . line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97898
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer


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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209044
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on
Authorized officer 	Authorized officer


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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64 . line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97899
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer

Form PCT/RO/134 (July 1992)


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Applicant's or agent's file reference number	PS001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209045</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

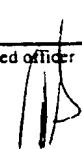
For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer

Applicant's or agent's file reference number	P001PCT	365	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64 line N/A	
B. IDENTIFICATION OF DEPOSIT	
Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97900
C. ADDITIONAL INDICATIONS (leave blank if not applicable)	
This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	


For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on
Authorized officer 	Authorized officer

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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209046
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	
For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on.
Authorized officer: 	Authorized officer


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Applicant's or agent's file reference number	S001PCT	International applicant	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 65, line N/A	
B. IDENTIFICATION OF DEPOSIT	
Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit April 28, 1997	Accession Number 209010
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	


For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer

Applicant's or agent's file reference number	S001PCT	368	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 65, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 29, 1997	Accession Number 209085
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	


For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on
Authorized officer 	Authorized officer

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Applicant's or agent's file reference number	'S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97897
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	
For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer


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Applicant's or agent's file reference number	PS001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 65 line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209043
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

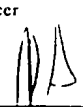
For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer

Applicant's or agent's file reference number	S001PCT	371	International applicant	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)


A. The indications made below relate to the microorganism referred to in the description on page <u>73</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution <p style="text-align: center;">American Type Culture Collection</p>	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit <u>September 4, 1997</u>	Accession Number <u>209236</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

<p>For receiving Office use only</p> <p><input checked="" type="checkbox"/> This sheet was received with the international application</p> <p>Authorized officer </p>	<p>For International Bureau use only</p> <p><input type="checkbox"/> This sheet was received by the International Bureau on</p> <p>Authorized officer</p>
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Applicant's or agent's file reference number	S001PCT	372	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 73, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 29, 1997	Accession Number 209084
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	
For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on
Authorized officer 	Authorized officer

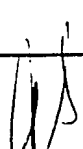
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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 76 line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209048
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	


For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer

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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 76, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97902
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit") 	
For receiving Office use only <input checked="" type="checkbox"/> This sheet was received with the international application Authorized officer 	For International Bureau use only <input type="checkbox"/> This sheet was received by the International Bureau on Authorized officer


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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 77, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97903
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

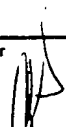
For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on
Authorized officer 	Authorized officer

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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>77</u> . line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209049
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	
For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on
Authorized officer 	Authorized officer


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Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 80 line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97904
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
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378

Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 80 line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209050
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit") 	

For receiving Office use only <input checked="checked" type="checkbox"/> This sheet was received with the international application Authorized officer 	For International Bureau use only <input type="checkbox"/> This sheet was received by the International Bureau on: Authorized officer
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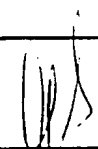
379

Applicant's or agent's file reference number	S001PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>82</u> line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit April 4, 1997	Accession Number 97976
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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380

Applicant's or agent's file reference number	5001PCT	International applicant	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 64, line N/A	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209047
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
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What Is Claimed Is:

1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:
- 5 (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- 10 (c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- 15 (e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, having biological activity;
- 20 (f) a polynucleotide which is a variant of SEQ ID NO:X;
- (g) a polynucleotide which is an allelic variant of SEQ ID NO:X;
- (h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;
- (i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.
- 25
2. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.
- 30
3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.
- 35

4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

5

5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.

10

6. The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.

15

7. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.

8. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.

20

9. A recombinant host cell produced by the method of claim 8.

10. The recombinant host cell of claim 9 comprising vector sequences.

25

11. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:

(a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z, having biological activity;

30

(c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

35

(e) a secreted form of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(f) a full length protein of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

- (g) a variant of SEQ ID NO:Y;
- (h) an allelic variant of SEQ ID NO:Y; or
- (i) a species homologue of the SEQ ID NO:Y.

5 12. The isolated polypeptide of claim 11, wherein the secreted form or the full length protein comprises sequential amino acid deletions from either the C-terminus or the N-terminus.

10 13. An isolated antibody that binds specifically to the isolated polypeptide of claim 11.

 14. A recombinant host cell that expresses the isolated polypeptide of claim 11.

15 15. A method of making an isolated polypeptide comprising:
 (a) culturing the recombinant host cell of claim 14 under conditions such that said polypeptide is expressed; and
 (b) recovering said polypeptide.

20 16. The polypeptide produced by claim 15.

 17. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 11 or the polynucleotide of claim 1.

25 18. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
 (a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and
 (b) diagnosing a pathological condition or a susceptibility to a pathological
30 condition based on the presence or absence of said mutation.

 19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
 (a) determining the presence or amount of expression of the polypeptide of
35 claim 11 in a biological sample; and
 (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

20. A method for identifying a binding partner to the polypeptide of claim 11 comprising:

- 5 (a) contacting the polypeptide of claim 11 with a binding partner; and
(b) determining whether the binding partner effects an activity of the polypeptide.

21. The gene corresponding to the cDNA sequence of SEQ ID NO:Y.

10 22. A method of identifying an activity in a biological assay, wherein the method comprises:

- (a) expressing SEQ ID NO:X in a cell;
(b) isolating the supernatant;
(c) detecting an activity in a biological assay; and
15 (d) identifying the protein in the supernatant having the activity.

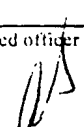
23. The product produced by the method of claim 22.

Applicant's or agent's file reference number	PS0017	International application	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>64</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>February 26, 1997</u>	Accession Number <u>97900</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	01PCT	International application number	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209043</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PS001	International application No.	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>64</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209044</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file
reference number

01PCT

International appl

o. Unassigned

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209045</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
in respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).	
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

Page 2

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

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NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PS0018	International application No.	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>64</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209046</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit") <div style="height: 100px;"></div>	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

Page 2

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	01PCT	International application No. Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>64</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209047</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g. "Accession Number of Deposit")	

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	PS001B	International application No.	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>76</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209048</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

Page 2

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	01PCT	International app	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>77</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209049</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

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UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	PS001P	International application	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>80</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 15, 1997</u>	Accession Number <u>209050</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>in respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

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NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	001PCT	International application no.	unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>73</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit September 4, 1997	Accession Number 209236 ..
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications. e.g., "Accession Number of Deposit")	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer: </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer: </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	PS001	International application No.	Assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>April 28, 1997</u>	Accession Number <u>209010</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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CANADA

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NORWAY

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AUSTRALIA

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FINLAND

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Page 2

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	001PCT	International app. No.	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution <i>(including postal code and country)</i> <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>May 29, 1997</u>	Accession Number <u>209085</u>
C. ADDITIONAL INDICATIONS <i>(leave blank if not applicable)</i> This information is continued on an additional sheet <input type="checkbox"/>	
<p>in respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE <i>(if the indications are not for all designated States)</i>	
E. SEPARATE FURNISHING OF INDICATIONS <i>(leave blank if not applicable)</i>	
The indications listed below will be submitted to the International Bureau later <i>(specify the general nature of the indications, e.g., "Accession Number of Deposit")</i>	

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file
reference number

PS001

International applicatio

to assigned

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description
on page 64, line N/A

B. IDENTIFICATION OF DEPOSIT

Further deposits are identified on an additional sheet ☒

Name of depositary institution American Type Culture Collection

Address of depositary institution (including postal code and country)

12301 Parklawn Drive
Rockville, Maryland 20852
United States of America

Date of deposit February 26, 1997

Accession Number 97901 ..

C. ADDITIONAL INDICATIONS (leave blank if not applicable)

This information is continued on an additional sheet ☐

In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).

D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)

E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)

The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

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UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	01PCT	International application No. Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>77</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>February 26, 1997</u>	Accession Number <u>97903</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
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CANADA

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NORWAY

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Page 2

UNITED KINGDOM

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DENMARK

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SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PS001	International application No.	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>64</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97898
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	001PCT	International application No. Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>80</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>February 26, 1997</u>	Accession Number <u>97904</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit") <div style="height: 100px;"></div>	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> Authorized officer </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

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DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

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NETHERLANDS

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Applicant's or agent's file
reference number

PS001P

International application No. assigned

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description
on page 73, line N/A

B. IDENTIFICATION OF DEPOSIT

Further deposits are identified on an additional sheet ☒

Name of depositary institution American Type Culture Collection

Address of depositary institution (including postal code and country)

12301 Parklawn Drive
Rockville, Maryland 20852
United States of America

Date of deposit May 29, 1997

Accession Number 209084

C. ADDITIONAL INDICATIONS (leave blank if not applicable)

This information is continued on an additional sheet ☐

In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).

D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)

E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)

The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	01PCT	International application No. Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>64</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>February 26, 1997</u>	Accession Number <u>97899</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

Page 2

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

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NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PS001	International application	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>65</u> . line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>February 26, 1997</u>	Accession Number <u>97897</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

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AUSTRALIA

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FINLAND

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Page 2

UNITED KINGDOM

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DENMARK

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SWEDEN

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NETHERLANDS

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Applicant's or agent's file reference number	901PCT	International app	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>82</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>April 4, 1997</u>	Accession Number <u>97976</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

<p style="text-align: center;">For receiving Office use only</p> <div style="border: 1px solid black; padding: 5px;"> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div style="border: 1px solid black; padding: 5px; height: 40px;"> Authorized officer </div>	<p style="text-align: center;">For International Bureau use only</p> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; height: 40px;"> Authorized officer </div>
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CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

Page 2

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

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AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

Page 2

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

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NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	P/S001	International application	assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>76</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>12301 Parklawn Drive</u> <u>Rockville, Maryland 20852</u> <u>United States of America</u>	
Date of deposit <u>February 26, 1997</u>	Accession Number <u>97902</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28 (4) EPC).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

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225 X 2.



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C12N 15/12, 5/10, 1/21, C07K 14/47, 16/18, C12Q 1/68, G01N 33/50, 33/53, 33/68, A61K 38/17	A3	(11) International Publication Number: WO 98/39446 (43) International Publication Date: 11 September 1998 (11.09.98)																														
(21) International Application Number: PCT/US98/04482 (22) International Filing Date: 6 March 1998 (06.03.98) (30) Priority Data: <table border="0" style="width: 100%;"> <tr> <td style="width: 40%;">60/040,162</td> <td style="width: 40%;">7 March 1997 (07.03.97)</td> <td style="width: 20%;">US</td> </tr> <tr> <td>60/040,333</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/038,621</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/040,161</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/040,626</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/040,334</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/040,336</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/040,163</td> <td>7 March 1997 (07.03.97)</td> <td>US</td> </tr> <tr> <td>60/043,580</td> <td>11 April 1997 (11.04.97)</td> <td>US</td> </tr> <tr> <td>60/043,568</td> <td>11 April 1997 (11.04.97)</td> <td>US</td> </tr> </table> <p><i>(Continued on the following page)</i></p> (71) Applicant (for all designated States except US): HUMAN GENOME SCIENCES, INC. [US/US]; 9410 Key West Avenue, Rockville, MD 20850 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): RUBEN, Steven, M. [US/US]; 18528 Heritage Hills Drive, Olney, MD 20832 (US). ROSEN, Craig, A. [US/US]; 22400 Rolling Hill Road, Laytonsville, MD 20882 (US). FISCHER, Carrie, L. [US/US]; 5810 Hall Street, Burke, VA 22015 (US). SOPER, Daniel, R. [US/US]; 15050 Stillfield Place, Centreville, VA 22020 (US). CARTER, Kenneth, C. [US/US]; 11601 Brandy Hall Lane, North Potomac, MD 20878 (US). BEDNARIK, Daniel, P. [US/US]; 8822 Blue Sea Drive, Columbia, MD 21046 (US). ENDRESS, Gregory, A. [US/US]; 9729 Clagett Farm Drive, Potomac, MD 20854 (US). YU, Guo-Liang [CN/US]; 13524 Straw Bale Lane, Darnestown, MD 20878 (US). NI, Jian [CN/US]; 5502 Manorfield Road, Rockville, MD 20853 (US). FENG, Ping [CN/US]; 4 Relda Court, Gaithersburg, MD 20878 (US). YOUNG, Paul, E. [US/US]; 122 Beckwith Street, Gaithersburg, MD 20878 (US). GREENE, John, M. [US/US]; 872 Diamond Drive, Gaithersburg, MD 20878 (US). FERRIE, Ann, M. [US/US]; 13203 L Astoria Hill Court, Germantown, MD 20874 (US). DUAN, Roxanne [US/US]; 4541 Fairfield Drive, Bethesda,		60/040,162	7 March 1997 (07.03.97)	US	60/040,333	7 March 1997 (07.03.97)	US	60/038,621	7 March 1997 (07.03.97)	US	60/040,161	7 March 1997 (07.03.97)	US	60/040,626	7 March 1997 (07.03.97)	US	60/040,334	7 March 1997 (07.03.97)	US	60/040,336	7 March 1997 (07.03.97)	US	60/040,163	7 March 1997 (07.03.97)	US	60/043,580	11 April 1997 (11.04.97)	US	60/043,568	11 April 1997 (11.04.97)	US	MD 20814 (US). HU, Jing-Shan [CN/US]; 1247 Lakeside Drive #3034, Sunnyvale, CA 94086 (US). GRAVES, Kimberly, A. [US/US]; 12805 Atlantic Avenue, Rockville, MD 20851 (US). OLSEN, Henrik, S. [DK/US]; 182 Kendrick Place #24, Gaithersburg, MD 20878 (US). EBNER, Reinhard [DE/US]; 9906 Shelburne Terrace #316, Gaithersburg, MD 20878 (US). BREWER, Laurie, A. [US/US]; 14920 Mt. Nebo Road, Poolesville, MD 20837 (US). MOORE, Paul, A. [GB/US]; Apartment #104, 1908 Holly Ridge Drive, McLean, VA 22102 (US). SHI, Yanggu [CN/US]; 437 West Side Drive, Gaithersburg, MD 20878 (US). LAFLEUR, David, W. [US/US]; 1615 Q Street, N.W. #807, Washington, DC 20009 (US). LI, Yi [CN/US]; 1247 Lakeside Drive #3034, Sunnyvale, CA 94086 (US). ZENG, Zhizhen [CN/US]; 13950 Saddleview Drive, Gaithersburg, MD 20878 (US). KYAW, Hla [BU/US]; 520 Sugarbush Circle, Frederick, MD 21703 (US). (74) Agents: BROOKES, Anders, A. et al.; Human Genome Sciences, Inc., 9410 Key West Avenue, Rockville, MD 10850 (US). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>With an indication in relation to a deposited microorganism furnished under Rule 13^{bis} separately from the description.</i> <i>Date of receipt by the International Bureau:</i> 06 April 1998 (06.04.98) (88) Date of publication of the international search report: 23 December 1998 (23.12.98)
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60/040,333	7 March 1997 (07.03.97)	US																														
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60/040,161	7 March 1997 (07.03.97)	US																														
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60/043,580	11 April 1997 (11.04.97)	US																														
60/043,568	11 April 1997 (11.04.97)	US																														
(54) Title: 70 HUMAN SECRETED PROTEINS (57) Abstract <p>The present invention relates to 70 novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins.</p>																																

60/043,314	11 April 1997 (11.04.97)	US	60/047,598	23 May 1997 (23.05.97)	US	60/056,882	22 August 1997 (22.08.97)	US
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60/047,492	23 May 1997 (23.05.97)	US	60/056,872	22 August 1997 (22.08.97)	US			

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DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

INTERNATIONAL SEARCH REPORT

International Application No

PC 98/04482

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/12 C12N5/10 C12N1/21 C07K14/47 C07K16/18
C12Q1/68 G01N33/50 G01N33/53 G01N33/68 A61K38/17

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K C12Q G01N A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	L. HILLIER ET AL.: "The WashU-Merck EST Project 1997" EMBL SEQUENCE DATABASE, 6 March 1997, HEIDELBERG, FRG, XP002068123 zr78g10.r1 Soares NhHMPu S1 Homo sapiens cDNA clone 669570 5' similar to SW:FUCO_RAT P17164 Alpha-L-fucosidase precursor; Accession. Accession no. AA234924; --- -/--	1-3, 7-10,21

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

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- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

16 June 1998

Date of mailing of the international search report

16. 09. 1998

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HORNIG H.

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 98/04482

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	L. HILLIER ET AL.: "The WashU-Merck EST Project" EMBL SEQUENCE DATABASE, 15 December 1996, HEIDELBERG, FRG, XP002068124 z140b11.r1 Soares pregnant uterus NbHPU Homo sapiens cDNA clone 504381 5' similar to TR:G182779 Lysosomal Enzyme Alpha-L-Fucosidase Accession no. AA151194	1-3, 7-10,21
X	--- L. HILLIER ET AL.: "The WashU-Merck EST Project" EMBL SEQUENCE DATABASE, 4 June 1996, HEIDELBERG, FRG, XP002068125 zc54a02.r1 Soares senescent fibroblasts NbHSF Homo sapiens cDNA clone 326090 5' similar to SW:FUCO_HUMAN P4066 tissue Alpha-L-Fucosidase precursor; Accession no. W52490	1-3, 7-10,21
A	--- WO 97 07198 A (GENETICS INSTITUT) 27 February 1997 see the whole document	1-23
A	--- WO 97 04097 A (GENETICS INST) 6 February 1997 see the whole document	1-23
A	--- US 5 536 637 A (JACOBS KENNETH) 16 July 1996 see the whole document	1-23
A	--- JACOBS K ET AL: "A novel method for isolating eukaryotic cDNA clones encoding secreted proteins." KEYSTONE SYMPOSIUM ON DENDRITIC CELLS: ANTIGEN PRESENTING CELLS OF T AND B LYMPHOCYTES, TAOS, NEW MEXICO, USA, MARCH 10-16, 1995. JOURNAL OF CELLULAR BIOCHEMISTRY SUPPLEMENT 0 (21A). 1995. 19. ISSN: 0733-1959, XP002027246 abstract no. C1-207 see abstract	1-23
A	--- WO 90 14432 A (GENETICS INST) 29 November 1990 see the whole document	1-23
A	--- WO 96 17925 A (IMMUNEX CORP) 13 June 1996 see the whole document	1-23

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INTERNATIONAL SEARCH REPORT

International Application No

PC 98/04482

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>T. OCCHIODORO ET AL.: "Human alpha-L-Fucosidase: Complete coding sequence from cDNA clones" BIOCHEM. AND BIOPHYS. RES. COMMUNICATIONS, vol. 164, no. 1, 16 October 1989, ACADEMIC PRESS, NEW YORK, US, pages 439-445, XP002068126 cited in the application see the whole document -----</p>	1-23

INTERNATIONAL SEARCH REPORT

mational application No.

PCT/US 98/04482

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claim 17 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see further information sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

see further information sheet

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

Invention 1. Claims: (1-23) partially

-An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence consisting of SEQ ID no. 11; wherein said polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein encoding the sequence of SEQ ID no. 134 or the polypeptide encoded by the cDNA sequence included in ATCC Deposit no: HGCMD20, which is hybridizable to SEQ ID no.11; a recombinant vector comprising said isolated nucleic acid molecule; a method of making a recombinant host cell comprising said isolated nucleic acid molecule; a recombinant host cell comprising said vector; an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence consisting of SEQ ID no. 134; an isolated antibody that binds specifically to said isolated polypeptide; a recombinant host cell that expresses said isolated polypeptide; a method of making said polypeptide; a method for preventing, treating or ameliorating a medical condition which comprises administering to a mammalian subject a therapeutically effective amount of said polypeptide; a method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject using said polynucleotide and/or polypeptide sequences; a method for identifying a binding partner to said polypeptide; a gene corresponding to the cDNA sequences of SEQ ID no.11; a method for identifying an activity in a biological assay, by using the expression of SEQ ID no. 134;

Inventions 2 to 70. Claims: (1-23) partially

-Idem as subject 1 but limited to gene nos. 2 to 70 respectively cDNA clone sequences HLDBG33 to HMCAB89.
(Invention 2 is limited to SEQ ID nos.12,81,135, and 204;
Invention 3 is limited to SEQ ID nos.13 and 136;;
Invention 70 is limited to SEQ ID nos.80 and 203;)

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/04482

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9707198 A	27-02-97	US 5707829 A AU 6712396 A AU 6768596 A EP 0839196 A EP 0851875 A WO 9704097 A	13-01-98 18-02-97 12-03-97 06-05-98 08-07-98 06-02-97
WO 9704097 A	06-02-97	US 5707829 A AU 6712396 A EP 0839196 A AU 6768596 A EP 0851875 A WO 9707198 A	13-01-98 18-02-97 06-05-98 12-03-97 08-07-98 27-02-97
US 5536637 A	16-07-96	US 5712116 A	27-01-98
WO 9014432 A	29-11-90	US 5580753 A AT 147436 T AU 637620 B AU 5928990 A CA 2056997 A DE 69029657 D DK 473724 T EP 0473724 A ES 2099096 T JP 4506006 T US 5734037 A US 5414071 A	03-12-96 15-01-97 03-06-93 18-12-90 24-11-90 20-02-97 14-04-97 11-03-92 16-05-97 22-10-92 31-03-98 09-05-95
WO 9617925 A	13-06-96	AU 4639396 A CA 2206488 A FI 972390 A NO 972455 A	26-06-96 13-06-96 05-06-97 06-08-97